

FORM PTO-1390 (REV. 11-2000) <b>TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371</b>		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTORNEY'S DOCKET NUMBER 018422-000310US
		U.S. APPLICATION NO. (If known, see 37 CFR 1.5) <b>09/857378</b>	
INTERNATIONAL APPLICATION NO. PCT/US99/28666	INTERNATIONAL FILING DATE December 3, 1999	PRIORITY DATE CLAIMED December 4, 1998	
<b>TITLE OF INVENTION</b> METHOD FOR THE IMMOBILIZATION OF OLIGONUCLEOTIDES			
<b>APPLICANT(S) FOR DO/EO/US</b> CHRISTOPHER P. ADAMS, JOSEPH D. KITTLE			
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:			
<p>1. <input checked="" type="checkbox"/> This is a <b>FIRST</b> submission of items concerning a filing under 35 U.S.C. 371.</p> <p>2. <input type="checkbox"/> This is a <b>SECOND</b> or <b>SUBSEQUENT</b> submission of items concerning a filing under 36 U.S.C. 371.</p> <p>3. <input checked="" type="checkbox"/> This is an express request to begin national examination procedures (35 U.S.C. 371(f)). The submission must include items (5), (6), (9) and (21) indicated below.</p> <p>4. <input checked="" type="checkbox"/> The US has been elected by the expiration of 19 months from the priority date (Article 31).</p> <p>5. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 37(c)(2))</p> <ol style="list-style-type: none"> <li><input type="checkbox"/> is attached hereto (required only if not communicated by the International Bureau).</li> <li><input type="checkbox"/> has been communicated by the International Bureau</li> <li><input checked="" type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US).</li> </ol> <p>6. <input type="checkbox"/> An English language translation of the International Application as filed (35 U.S.C. 371(c)(2)).</p> <ol style="list-style-type: none"> <li><input type="checkbox"/> is attached hereto.</li> <li><input type="checkbox"/> has been previously submitted under 35 U.S.C. 154(d)(4).</li> </ol> <p>7. <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3)).</p> <ol style="list-style-type: none"> <li><input type="checkbox"/> are attached hereto (required only if not communicated by the International Bureau).</li> <li><input type="checkbox"/> have been communicated by the International Bureau.</li> <li><input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired.</li> <li><input checked="" type="checkbox"/> have not been made and will not be made.</li> </ol> <p>8. <input type="checkbox"/> An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).</p> <p>9. <input type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).</p> <p>10. <input type="checkbox"/> An English language translation of the annexes of the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).</p> <p><b>Items 11 to 20 below concern document(s) or information included:</b></p> <p>11. <input type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98.</p> <p>12. <input type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.</p> <p>13. <input type="checkbox"/> A FIRST preliminary amendment.</p> <p>14. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment.</p> <p>15. <input type="checkbox"/> A substitute specification.</p> <p>16. <input type="checkbox"/> A change of power of attorney and/or address letter.</p> <p>17. <input type="checkbox"/> A computer-readable form of the sequence listing in accordance with PCT Rule 13ter.2 and 35 U.S.C. 1.821 – 1.825.</p> <p>18. <input type="checkbox"/> A second copy of the published international application under 36 U.S.C.</p> <p>19. <input type="checkbox"/> A second copy of the English language translation of the international application under 35 U.S.C. 154(d)(4).</p> <p>20. <input type="checkbox"/> Other items or information:</p>			

I/S Application no. (if known) see 37 CFR 1.57 <b>09/857378</b>	INTERNATIONAL APPLICATION NO. PCT/US99/28666	ATTORNEY'S DOCKET NUMBER 018422-000310US			
21. <input checked="" type="checkbox"/> The following fees are submitted:		CALCULATIONS PTO USE ONLY			
<b>BASIC NATIONAL FEE (37 CFR 1.492(A) (1) - (5)):</b>					
Neither international preliminary examination fee (37 CFR 1.492) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO .....		\$1000.00			
International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search report prepared by the EPO of JPO .....		\$860.00			
International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO .....		\$710.00			
International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4) .....		\$690.00			
International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)(4) .....		\$100.00			
<b>ENTER APPROPRIATE BASIC FEE AMOUNT =</b>		\$100			
Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(e)).		\$			
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE	\$	
Total claims	65 - 20 =	45	x \$18.00	\$810	
Independent claims	8 - 3 =	5	x \$80.00	\$400	
<b>MULTIPLE DEPENDENT CLAIM(S) (if applicable)</b>		+ 270.00	\$		
<b>TOTAL OF ABOVE CALCULATIONS =</b>		\$1,310			
<input checked="" type="checkbox"/> Applicant claims small entity status. See 37 CFR 1.27. The fees indicated above are reduced by 1/2.		+ \$655			
<b>SUBTOTAL =</b>		\$655			
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).		\$			
<b>TOTAL NATIONAL FEE =</b>		\$655			
Fee for recording the enclosed assignment (37 CFR 1.2(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property		+ \$			
<b>TOTAL FEES ENCLOSED =</b>		\$655			
		<b>Amount to be refunded:</b>	\$		
		<b>charged:</b>	\$		
a. <input type="checkbox"/> A check in the amount of \$ _____ to cover the above fees is enclosed.					
b. <input checked="" type="checkbox"/> Please charge my Deposit Account No. <u>20-1430</u> in the amount of <u>\$655</u> to cover the above fees.					
c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. <u>20-1430</u> . A duplicate copy of this sheet is enclosed.					
d. <input type="checkbox"/> Fees are to be charged to a credit card. <b>WARNING:</b> Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038.					
<b>NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b) must be filed and granted to restore the application to pending status.</b>					
SEND ALL CORRESPONDENCE TO:					
Eugenia Garrett-Wackowski Townsend and Townsend and Crew LLP Two Embarcadero Center, 8th fl. San Francisco, CA 94111					
Annette S. Parent NAME					
42,058 REGISTRATION NUMBER					

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Rec'd PCT/PTO 04 JUN 2001

"METHOD FOR THE IMMOBILIZATION OF OLIGONUCLEOTIDES"

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**CROSS-REFERENCES TO RELATED APPLICATIONS**

This application claims priority to U.S. Provisional Application Serial No. 60/110,891, filed on December 4, 1998. The disclosure of the Provisional Application is incorporated herein by reference in its entirety for all purposes.

10

**BACKGROUND OF THE INVENTION**

A perennial goal in the pharmacological arts is the development of methods and compositions to facilitate the specific delivery of therapeutic and other agents to cells and tissues that would benefit from such treatment, and the avoidance of the general physiological effects of the inappropriate delivery of such agents to other cells or tissues of the body. Recently, the advent of recombinant DNA technology and genetic engineering has provided the pharmacological arts with a wide spectrum of new agents that are functional genes carried in recombinant expression constructs capable of mediating expression of these genes in host cells. These developments have carried the promise of "molecular medicine," specifically gene therapy, whereby a defective gene is replaced by an exogenous copy of its cognate, functional gene, thereby alleviating a variety of genetic diseases.

An ever-expanding array of genes, the abnormal expression of which, is associated with life-threatening human diseases is being cloned and identified. The ability to express such cloned genes in humans will ultimately permit the prevention and/or cure of many important human diseases, which now are either poorly treated or are untreatable by available therapies. As an example, *in vivo* expression of cholesterol-regulating genes, genes which selectively block the replication of HIV, or tumor-suppressing genes in human patients should dramatically improve treatment of heart disease, HIV, and cancer, respectively. However, currently available gene delivery strategies have been unable to produce a high level of generalized transgene expression *in vivo* after systemic administration to a mammalian host. This inability has precluded the development of effective gene therapy for most human diseases.

The various approaches to gene therapy include both different goals and different means of achieving those goals. The goals include gene replacement, gene correction and gene augmentation. In gene replacement, a mutant gene sequence is specifically removed from the genome and replaced with a normal, functional gene. In gene correction, a mutant gene sequence is corrected without any additional changes in the target genome. In gene augmentation, the expression of mutant genes in defective cells is modified by introducing foreign normal genetic sequences.

The means to reach the above goals include "*ex vivo*" transfection of a target cell, followed by introduction of the transformed cells into a suitable organ in the host mammal. *Ex vivo* techniques include transfection of cells *in vitro* with either naked DNA or DNA encapsulated in, for example, liposomes, followed by introduction into a host organ ("*ex vivo*" gene therapy).

There are several drawbacks to *ex vivo* therapy. For example, if only differentiated, replicating cells are infected, the newly introduced gene function will be lost as those cells mature and die. *Ex vivo* approaches can be used to transfect only a limited number of cells and cannot be used to transfect cells which are not first removed from the body. The above methods involve integration of new genetic material into the cell genome and thus constitute permanent changes to the host genome. However, some gene augmentation can be achieved using methods that do not involve changes to the genome, but which introduce DNA into a host cell where it is maintained primarily in an extrachromosomal or episomal form.

The greatest drawback to the achievement of effective gene therapy has been the inability in the art to introduce recombinant expression constructs encoding functional eukaryotic genes into cells and tissues *in vivo*. While it has been recognized as desirable to increase the efficiency and specificity of administration of gene therapy agents to the cells of the relevant tissues, the goal of specific delivery has not yet been achieved.

In methods other than *ex vivo* methods, genetic material is transferred into target cells without the use of vectors or carriers. For example, genetic material is introduced systemically through an intravenous or intraperitoneal injection for *in vivo* applications, or it is introduced to the site of action by direct injection into that area. For example, it has long been recognized that DNA, by itself, injected into various tissues, will enter cells and produce a protein eliciting an immune response. See, e.g., Atanasiu *et al.*, *Academie des Sciences (Paris)* 254:4228-30 (1962); Israel *et al.*, *J. Virol.* 29:990-96

(1979); Will *et al.*, *Nature*, **299**:740-42 (1982); Robinson, WO 86/00930, published 13 Feb. 1986; Felgner *et al.*, WO 90/11092, published 4 Oct. 1990; and Debs *et al.*, WO 93/24640, published 9 Dec. 1993. DNA by itself, however, is hydrophilic and the hydrophobic character of the cellular membrane poses a significant barrier to the transfer 5 of naked DNA across it. Accordingly, it is generally preferred to use facilitators that enhance the transfer of DNA into cells on direct injection.

Facilitators that have been used are generally polycationic in nature. For example, polylysine has been widely investigated as a facilitator (Soeda *et al.*, *Gene Ther.* **5**:1410 (1998)). Poly-lysine is not unique in providing a polycationic framework for the entry of DNA into cells. DEAE-dextran has also been shown to be effective in promoting RNA and DNA entry into cells; *see*, Juliano *et al.*, *Exp. Cell. Res.* **73**:3-12 (1972); and Mayhew *et al.*, *Exp. Cell. Res.* **77**:409-414 (1973). More recently, a dendritic cascade copolymer of ethylenediamine and methyl acrylate has been shown to be useful in providing a carrier of DNA which facilitates entry into cells; *see*, Haensler *et al.*, *Bioconj. Chem.* **4**:372-379 (1993). An alkylated polyvinylpyridine polymer has also been used to facilitate DNA entry into cells; *see*, Kabanov, *et al.*, *Bioconj. Chem.* **4**:448-454 (1993). None of these references suggests forming a polymer from subunits including both an ethylene moiety and a nucleic acid.

Many polycationic facilitators or carriers are used to reversibly complex a polyanionic nucleic acid by an ionic binding mechanism. For example, Hennick *et al.*, WO 97/15680, published 1 May 1997, have used a synthetic transfection system comprising a water dispersible or water soluble carrier formed from polyacrylate, polyacrylamide and derivatives thereof. The carriers are substituted with cationic groups, which reversibly complex a nucleic acid. The use of cationic carriers to reversibly complex therapeutic nucleic acids is limited in its scope; when uncharged nucleic acid derivatives (*e.g.*, phosphorothioates) are utilized, an ionic bonding mechanism will not be operative. Other facilitators take advantage of the ability of nucleic acids to form hydrogen bonds. Mumper *et al.* (*Pharm. Res.* **13**:701 (1996)) have reported the use of polyvinyl derivatives (*e.g.*, polyvinylpyrrolidone, polyvinylalcohol) as interactive polymers for controlled gene delivery to muscle. The polyvinyl derivatives bind reversibly to the nucleic acid via a hydrogen bonding mechanism. There is no suggestion in either Hennick *et al.* or Mumper *et al.* to form polymers with nucleic acids derivatized with ethylene-containing moieties or that a nucleic acid can be polymerized to form a conjugate to which the nucleic acid is covalently bound.

Nucleic acids that are substituted with an olefin are known in the art. For example, Nagatsuge *et al.*, (*Tetrahedron* 9:3035 (1997)) have described the synthesis of 2-aminopurine derivatives with a C<sup>6</sup>-substituted olefin. These agents are used to form crosslinks with a target nucleobase due to their proximity to the base in a sense-antisense duplex. There is no suggestion in this reference to incorporate the olefinic group of the olefinically derivatized base into the backbone of a polymer. Furthermore, Mosaic Technologies, Inc. has introduced ethylene-containing phosphoramidite linkers that allow an ethylene group to be tethered to a nucleic acid (Acrydite<sup>TM</sup>). The ethylene-derivatized nucleic acids are generally immobilized by polymerization in an acrylamide gel. See, for example, [www.mostek.com](http://www.mostek.com); Kenney *et al.*, *Biotechniques* 25:516 (1998). The ethylene-linked nucleic acids have not been suggested for incorporation into particles or substantially water-soluble polymers. Moreover, they have not been suggested as components of a vehicle for delivering nucleic acids to a target cell.

In spite of active research in the area of delivery vehicles for therapeutic nucleic acids in gene therapy, an agent has not been developed that allows polymeric carriers to be made from ethylene-derivatized nucleic acids. In view of the number of polymerization and addition reaction pathways available to compounds bearing the ethylene group, a set of nucleic acid-derivatized polymer backbones having a great deal of structural diversity can quickly and easily be generated from nucleic acids bearing an ethylene group. The ability to rapidly generate compounds having a broad range of structural diversity allows for delivery vehicles having improved pharmacological properties to be quickly identified from a pool of diverse structures. The compounds with improved properties are useful in methods, such as gene therapy. Quite surprisingly, the present invention provides such compounds.

25

## SUMMARY OF THE INVENTION

It has now been discovered that nucleic acid polymers, monomers and nucleic acid analogues can be tethered to a scaffold (*e.g.*, a polymer, oligomer, *etc.*) via a species comprising an ethylene moiety (*e.g.*, acrylic- or vinyl- containing reactive group). The ethylene group is generally tethered to the 3'- or 5'-hydroxyl of the nucleic acid or nucleic acid analogue, however, this group can be attached at any position of the nucleic acid chain and to any group on either the sugar or the base of the nucleic acid or nucleic acid analogue.

The incorporation into the nucleic acid of the ethylene derivatized group can be easily accomplished by utilizing a derivative such as a phosphoramidite to which the ethylene derivatized reactive group is tethered. Phosphoramidite and other appropriate nucleic acid chemistries are well known in the art and suitable reaction schemes will be apparent to those of skill in the art.

Thus, in a first aspect, the present invention provides a substantially water-soluble polymer including a first subunit comprising a first nucleic acid. The first subunit is incorporated into the polymer using a first subunit precursor. The first subunit precursor includes the first nucleic acid and an ethylene-containing moiety.

In a second aspect, the invention provides a polymeric particle including a first subunit. The first subunit includes a first nucleic acid. The first subunit is incorporated into the polymer using a first subunit precursor. The first subunit precursor includes an ethylene-containing moiety.

In a third aspect, the instant invention provides a pharmaceutical formulation including a pharmaceutically acceptable carrier and a substantially water-soluble polymer. The polymer includes a first subunit. The first subunit includes a first nucleic acid. The first subunit is incorporated into the polymer using a first subunit precursor including the first nucleic acid and an ethylene-containing moiety.

In a fourth aspect, the present invention provides a pharmaceutical formulation including a pharmaceutically acceptable carrier and a polymeric particle. The particle includes a first subunit. The first subunit includes a first nucleic acid. The first subunit is incorporated into the polymer using a first subunit precursor, which includes the first nucleic acid and an ethylene-containing moiety.

In a fifth aspect, the present invention provides a method for treating or preventing a condition in a subject. The method includes administering to the subject a substantially water-soluble polymer in an amount effective to treat or prevent the condition. The polymer comprises a first subunit, which includes a first nucleic acid. The first subunit is incorporated into the polymer using a first subunit precursor, which includes the first nucleic acid and an ethylene-containing moiety.

In a further aspect, the present invention provides a method for treating or preventing a condition in a subject. The method includes administering to the subject a polymeric particle in an amount effective to treat or prevent the condition. The particle includes a first subunit, which includes a first nucleic acid. The first subunit is

incorporated into the polymer using a first subunit precursor. The precursor includes the first nucleic acid and an ethylene-containing moiety.

In a still further aspect, the invention provides a method for introducing a polynucleotide into a eukaryotic cell in a living animal. The method includes contacting the cell with a composition comprising a substantially water-soluble polymer, which includes a first subunit. The first subunit includes a first nucleic acid. The first subunit is incorporated into the polymer using a first subunit precursor, which includes the first nucleic acid and an ethylene-containing moiety.

In another aspect, the invention provides a method for introducing a polynucleotide into a eukaryotic cell in a living animal. The method includes contacting the cell with a composition comprising a polymeric particle, which includes a first subunit. The first subunit includes a first nucleic acid. The first subunit is incorporated into the polymer using a first subunit precursor, which includes the nucleic acid and an ethylene-containing moiety.

Additional objects and advantages of the present invention will be apparent from the detailed description that follows.

## DETAILED DESCRIPTION OF THE INVENTION AND THE PREFERRED EMBODIMENTS

### A. Definitions

As used herein the term "polymer" refers to molecules having two or more subunits (e.g., dinucleotides).

As used herein, the term "nucleic acid" is used interchangeably with RNA and DNA and this term can refer to monomeric, oligomeric or polymeric species of these molecules. Moreover, nucleic acid analogues are incorporated within this definition.

"Alkyl" denotes straight-chain, branched-chain, saturated and unsaturated groups.

"Substituted alkyl" refers to alkyl as just described including one or more functional groups such as lower alkyl, aryl, acyl, halogen (i.e., alkylhalos, e.g., CF<sub>3</sub>), hydroxy, amino, alkoxy, alkylamino, acylamino, acyloxy, aryloxy, aryloxyalkyl, mercapto, both saturated and unsaturated cyclic hydrocarbons, heterocycles and the like. These groups may be attached to any carbon of the alkyl moiety. Moreover, the substitution can be pendent from the alkyl chain or interrupt the alkyl chain.

The term "aryl" is used herein to refer to an aromatic group, which may be a single aromatic ring or multiple aromatic rings which are fused together, linked covalently, or linked to a common group such as a methylene or ethylene moiety. The common linking group may also be a carbonyl as in benzophenone. The aromatic ring(s) 5 may include phenyl, naphthyl, biphenyl, diphenylmethyl and benzophenone among others. The term "aryl" encompasses "arylalkyl."

The term "arylalkyl" is used herein to refer to a subset of "aryl" in which the aryl group is attached to the another group of the compound by an alkyl group as defined herein.

10 "Substituted aryl" refers to aryl as just described including one or more functional groups such as lower alkyl, acyl, halogen, alkylhalos (e.g. CF<sub>3</sub>), hydroxy, amino, alkoxy, alkylamino, acylamino, acyloxy, mercapto and both saturated and unsaturated cyclic hydrocarbons which are fused to the aromatic ring(s), linked covalently or linked to a common group such as a methylene or ethylene moiety. The linking group 15 may also be a carbonyl such as in cyclohexyl phenyl ketone. The term "substituted aryl" encompasses "substituted arylalkyl."

"Substituted arylalkyl" defines a subset of "substituted aryl" wherein the substituted aryl group is attached to another group of the compound by an alkyl group as defined herein.

20 "Substituted" encompasses both single and multiple substitutions; the latter including multiple substitutions by the same substituent as well as mixtures of different substituents.

## B. Introduction

25 In accordance with the subject invention, nucleic acid constructs together with methods of preparation and use of these constructs are provided which allow for *in vivo* change of the genotype and/or modulation of the phenotype of cells in a plurality of tissues of a mammalian host, following introduction of the constructs into a circulating body fluid, organ or body cavity at a sufficient dose to cause transfection of tissues and/or 30 cells contacted by the nucleic acid. The tissues which are transformed include, for example, the lungs, heart, liver, bone marrow, spleen, lymph nodes, kidneys, thymus, skeletal muscle, ovary, uterus, stomach, small intestine, colon, pancreas, and brain in normal animals, as well as metastatic tumors and intravascular tumor emboli in tumor-bearing mammals. Particular cells which are transfected include, for example,

macrophages, alveolar type I and type II cells, hepatocytes, airway epithelial cells, vascular endothelial cells, cardiac myocytes, myeloblasts, erythroblasts, B-lymphocytes and T-lymphocytes. The circulating bodily fluid is generally blood, but intrathecal administration can also be used.

The invention provides compositions and pharmaceutical formulations including these compositions. Methods for using the compositions of the invention are also provided. Included are methods for treating or preventing a condition in a subject and methods for delivering genetic material into a cell.

10     1. *Compositions*

(a). **Substantially water-soluble polymers**

In a first aspect, the present invention provides a substantially water-soluble polymer comprising a first subunit, which includes a first nucleic acid. The first subunit is incorporated into the polymer using a first subunit precursor, which includes the first nucleic acid and an ethylene-containing moiety.

The polymers of the invention can have substantially any structure achievable by using subunits having a polymerizable or otherwise reactive ethylene moiety. Thus, the polymers can be homopolymers or copolymers of two or more structurally distinct subunits. The subunits themselves can be either monomeric or polymeric. Thus, it is within the scope of the present invention to construct a single polymer from two distinct polymer molecules having similar or quite different properties. Alternatively, one or more of the subunits can be a polymer to which monomeric subunits are appended. Other permutations of the conjugation of monomers and/or polymers will be apparent to those of skill in the art and are useful in preparing the compounds of the present invention.

In a preferred embodiment, the subunits of the present polymers are attached via a cleavable moiety. Many cleavable groups are known in the art. See, for example, Jung *et al.*, *Biochem. Biophys. Acta*, **761**:152-162 (1983); Joshi *et al.*, *J. Biol. Chem.* **265**:14518-14525 (1990); Zarling *et al.*, *J. Immunol.* **124**:913-920 (1980); Bouizar *et al.*, *Eur. J. Biochem.* **155**:141-147 (1986); Park *et al.*, *J. Biol. Chem.* **261**:205-210 (1986); Browning *et al.*, *J. Immunol.* **143**:1859-1867 (1989). Exemplary cleavable moieties can be cleaved using light, heat or reagents such as thiols, hydroxylamine, bases, periodate and the like.

Cleaveable groups preferred for use in the compounds of the invention include a cleaveable moiety that is a member selected from the group consisting of disulfide, ester, imide, carbonate, nitrobenzyl, phenacyl and benzoin groups.

In an exemplary embodiment, the polymer includes at least two units joined by a photocleavable group. The polymer is delivered to a tissue, for example, a tumor, and after some duration for a desired degree of uptake or distribution to occur, a high intensity light is focused on the tissue. The light is of sufficient energy to cleave the photocleavable bonds and disrupt the polymeric backbone, and release the nucleic acid.

Still further preferred cleavable moieties are those that undergo cleavage due to a naturally occurring biological process. Exemplary cleavable groups according to this motif include, for example, a cleavable linker that is sensitive to the slightly acidic pH of endocytotic vacuoles. In one embodiment, the polymer is delivered to a desired tissue and taken up by the cell via encapsulation in an endocytotic vacuole, where it is cleaved into smaller fragments by the acidic environment of the vacuole.

Representative groups cleaved by biological processes include, for example, disulfides, esters, phosphodiesters and combinations thereof. Disulfides are cleaved *in vivo* by reducing enzymes and small molecule thiol transfer reagents. Esters undergo hydrolysis and are also catalytically cleaved by esterases. Phosphodiesters are cleaved by nucleases.

A representative disulfide-containing polymer is prepared as follows. A first ethylene-containing subunit, to which a first nucleic acid is tethered, is cross-linked, via a disulfide-containing crosslinking agent, to a second ethylene-containing subunit, to which a second nucleic acid is optionally tethered. Useful crosslinking agents are those that include both a disulfide group and two or more groups reactive with the ethylene-containing moieties of the subunits. A representative agent is N,N'-bis(acryloyl)cystamine (Hansen, *Anal. Biochem.* 76:37(1976), commercially available from Sigma, St. Louis, MO).

A representative ester-containing polymer is prepared as follows. A first ethylene-containing subunit, to which a first nucleic acid is tethered, is cross-linked, via an ester-containing crosslinking agent, to a second ethylene-containing subunit, to which a second nucleic acid is optionally tethered. Useful crosslinking agents are those that include both an ester group and two or more groups reactive with the ethylene-containing moieties of the subunits. Representative agents include, the bis-acryloyl- and bis-

methacryloyl-poly(ethyleneglycol) agents commercially available from Shearwater Polymers (Huntsville, AL).

Other ester containing crosslinking agents are easily prepared in a few steps from commercially available starting materials, such as the active ester of acrylic acid, acrylic acid-N-hydroxysuccinimide. This agent is used to prepare bis-acryloyl ester derivatives of, for example, a wide variety of unsubstituted and internally substituted  $\alpha$ -,  $\omega$ -diols. Other combinations of cleavable cross-linking agents and subunits of use in preparing the polymers of the invention will be apparent to those of skill in the art.

In another preferred embodiment, the nucleic acid and the ethylene-containing moiety are joined by a linking group. Using such linking groups, the properties of the polymer can be controlled. Properties that are usefully controlled include, for example, hydrophobicity, hydrophilicity, surface-activity and the distance of the nucleic acid from the polymer backbone. For example, in a polymer including a backbone of largely aliphatic character, the nucleic acid can be attached to the polymer backbone via a poly(ethyleneglycol) to enhance the hydrophilicity of the polymer or to impart additional steric freedom to the nucleic acid. Numerous other combinations of linking groups and polymer backbones are accessible to those of skill in the art.

The hydrophilicity of the polymer can be enhanced by incorporating linking groups that include polar moieties such as amines, hydroxyls and polyhydroxyls. Representative examples include, but are not limited to, polylysine, polyethyleneimine, poly(ethyleneglycol) and poly(propyleneglycol). Suitable functionalization chemistries and strategies for these compounds are known in the art. See, for example, Dunn, R.L., *et al.*, Eds. POLYMERIC DRUGS AND DRUG DELIVERY SYSTEMS, ACS Symposium Series Vol. 469, American Chemical Society, Washington, D.C. 1991.

The hydrophobicity of the polymer can be modulated by using a hydrophobic linking group such as, for example, long chain diamines, long-chain thiols,  $\alpha$ ,  $\omega$ -amino acids, etc. Representative hydrophobic spacers include, but are not limited to, 1,6-hexanediamine, 1,8-octanediamine, 6-aminohexanoic acid and 8-aminoctanoic acid.

The polymer can also be made surface-active by using a linking group having surfactant properties. Compounds useful for this purpose include, for example, aminated or hydroxylated detergent molecules such as, for example, 1-aminododecanoic acid.

In another embodiment, the linking group serves to distance the nucleic acid from the polymer backbone. Groups with this characteristic have several uses. For example, a nucleic acid held too closely to the polymer backbone may not be able to hybridize with an incoming complementary nucleic acid strand, or it may hybridize unacceptably slowly. When an incoming nucleic acid strand is itself sterically demanding, the hybridization can be undesirably slowed, or not occur at all, due to the monolithic polymer backbone hindering the approach of the two complementary strands. Thus, linking groups that provide distance between the nucleic acid and the polymer backbone can serve to enhance nucleic acid hybridization.

In a still further preferred embodiment, the linking group includes a cleavable moiety. The discussion above concerning cleavable moieties is generally applicable to this embodiment as well.

In another embodiment, the physicochemical characteristics (e.g., hydrophobicity, hydrophilicity, surface activity, conformation) of the polymer are altered by attaching a monovalent moiety which is different in composition than the constituents of the bulk polymer and which does not bear a nucleic acid. As used herein, "monovalent moiety" refers to organic molecules with only one reactive functional group. This functional group attaches the molecule to the polymer backbone. "Monovalent moieties" are to be contrasted with the bifunctional linking groups described above. Such monovalent groups are used to modify the hydrophilicity, hydrophobicity, binding characteristics, etc. of the polymer. Examples of groups useful for this purpose include long chain alcohols, amines, fatty acids, fatty acid derivatives, poly(ethyleneglycol) monomethyl ethers, etc.

In the polymers of the invention, the ethylene-containing moiety can have substantially any structure deemed useful for a particular application. Thus, the ethylene-containing moiety can be, for example, a saturated alkyl group an unsaturated alkyl group, a carbohydrate, an amino acid or peptide, a polyether, a polyamine, etc.

In a presently preferred embodiment, the ethylene-containing moiety includes a member selected from  $\text{CH}_2=\text{CHX}^1$ ,  $\text{CH}_2=\text{CX}^2\text{Y}^1$  and combinations thereof. In these ethylene-containing moieties  $X^1$ ,  $X^2$  and  $Y^1$  are members independently selected from H, (=O), NR<sup>1</sup>R<sup>2</sup>, OH, and OR<sup>3</sup>. R<sup>1</sup>, R<sup>2</sup> and R<sup>3</sup> are members independently selected from H, alkyl, substituted alkyl, aryl and substituted aryl.

In a still further preferred embodiment, R<sup>1</sup>, R<sup>2</sup> and R<sup>3</sup> are independently selected from H, alkyl and substituted alkyl, more preferably from H, alkyl and alkyl substituted with at least one moiety selected from OH, O and combinations thereof.

5 In yet another preferred embodiment, at least one of R<sup>1</sup>, R<sup>2</sup> and R<sup>3</sup> includes a moiety selected from poly(ethyleneglycol), poly(propyleneglycol) and combinations thereof.

In another preferred embodiment, the polymer backbone includes a group that is derived from a member selected from acrylate, acrylamide, C<sub>1</sub>-C<sub>6</sub> alkylacrylate, (alkyl)acrylamide, methylmethacrylate, triethyleneglycolmethacrylate, 10 poly(ethyleneglycol)methacrylate, hydroxyethylmethacrylate, glycerylmethacrylate, vinyl alcohol, ethylcyanoacrylate and combinations thereof.

**(i). Framework components**

In another preferred embodiment, the monomeric or polymeric ethylene-containing subunits are attached to a secondary polymeric component that is derivatized to allow for such attachment. In this embodiment, the polymeric moiety to which the subunits are attached is referred to as a "framework component." Examples of framework components suitable for use in the methods of the present invention include, but are not limited to, polymers, liposomes, micelles, colloids, biological particles and 15 non-biological particles (*e.g.*, silica beads, polymeric beads, gels, *etc.*). These various types of framework components are discussed briefly below and in more detail further below under the description of covalent and noncovalent frameworks. The detailed descriptions of each of these framework components are provided under the headings of covalent and noncovalent framework components only for ease of discussion and should 20 not be construed as limiting the scope of useful framework structures. It is understood that the present invention is intended to encompass all types of frameworks capable of presenting functional groups that are reactive towards the ethylene-containing moiety of the subunits.

As noted above, examples of framework components suitable for use in 25 the methods of the present invention include, but are not limited to, polymers, liposomes, micelles, colloids, dendrimers and biological particles. The terms "polymer" and "polymeric" are art-recognized terms and, as used herein, include reference to a structural framework including repeating monomer units. The terms also include reference to

homopolymers and copolymers. Linear polymers, branched polymers and cross-linked polymers are also encompassed within the terms "polymer" and "polymeric."

The terms "liposome," "micelles," and "colloids" are art-recognized terms and, as used herein, these terms also include the derivatized versions, *e.g.*, liposome derivatives, cross-linked liposomes, *etc.*

The term "biological particle" includes reference to both covalent molecules, *e.g.*, sugars, proteins, lipid, small molecules, protein aggregates, and nucleic acids, and noncovalent particles, *e.g.*, modified cells (*e.g.*, which have been derivatized, modified chemically or transfected with an exogenous nucleic acid), or modified viruses, *e.g.* viral particles. The use of "biological particles" as framework components is distinguished from such particles as they occur in their natural state because the subject framework components are typically modified to present a nucleic acid.

In each of the embodiments discussed hereinbelow, it is generally understood that prior to their being reacted with the ethylene-containing nucleic acid conjugate, the covalent framework component is either "naturally" or "synthetically primed" for reaction with the ethylene group of the conjugate. "Naturally primed" polymers are those that can be coupled to the ethylene-containing moiety without prior derivitization of the polymers. "Synthetically primed" polymers are those that undergo additional modification prior to conjugation with the ethylene group.

In an exemplary embodiment, a framework component that bears amines on its surface is reacted with an activated ester (*e.g.*, N-hydroxysuccinimide) of a compound, such as acrylic acid, thereby converting the amines to an ethylene-containing group that can be reacted with the ethylene-containing subunits discussed above. In another embodiment, the surface of the amine-containing polymer is utilized in an addition reaction, such as the Michael addition, in which case the amine-containing polymer is naturally primed for reaction with the ethylene-containing moiety of the conjugate. Many other priming strategies and reaction sequences can be utilized to prepare an appropriately reactive framework component. Such reaction sequences are well known, and easily accessible, to those of skill in the art.

### 30 ***Covalent Framework Components***

In one embodiment, the monomeric units of a framework component are joined covalently. Exemplary covalent frameworks include, but are not limited to, cross-linked liposomes, biological particles (*e.g.*, sugars, proteins, peptides, lipids, or small

molecules) and polymeric materials (*see, e.g.*, Siraganian *et al.*, *Immunochem.* **12**:149-155 (1975); Wofsy *et al.*, *J. Immunol.* **121**:593-601 (1978); Barlocco *et al.*, *Farmac.* **48**:387-96 (1993); Castagnino *et al.*, *Jpn. Heart J.* **31**:845-55 (1990); Costa *et al.*, *Biochem. Pharmacol.* **34**:25-30 (1985); Dembo *et al.*, *J. Immunol.* **122**:518-28 (1979); Holliger *et al.*, *Proc. Natl. Acad. Sci. U. S. A.* **90**:6444-8 (1993); Piergentili *et al.*, *Farmaco* **49**:83-7 (1994); Portoghesi *et al.*, *J. Med Chem.* **34**:1292-6 (1991); Kizuka *et al.*, *J. Am. Chem. Soc.* **30**:722-6 (1987)).

In certain embodiments, proteins, *e.g.*, albumin, can be used as a framework component for presenting large numbers of groups (Roy *et al.*, *Can. J. Chem.* **68**:2045-2054 (1990)), thereby mimicking natural glycoprotein inhibitors.

In presently preferred embodiments, polymers are used as the framework component. Polymers are a versatile framework system (*see, e.g.*, Spaltenstein *et al.*, *J. Am. Chem. Soc.* **113**:686 (1991); Mammen *et al.*, *J. Med. Chem.* **38**:4179 (1995)). In a preferred embodiment, the ethylene-containing nucleic acids of the present invention are attached to a framework component comprising a polymeric backbone through a linker group.

Polymers can be purchased from commercial sources or, alternatively, they can be prepared using methods known to those of skill in the art (*See, e.g.*, Sandler, *et al.*, *POLYMER SYNTHESES*; Harcourt, Brace: Boston, 1994; Shalaby *et al.*, *J. Polymers of Biological and Biomedical Significance (ACS Symposium Series 540)*; American Chemical Society: Washington, DC, 1994). Moreover, polymeric frameworks are easily, rapidly and convergently synthesized (*see, e.g.*, Spaltenstein *et al.*, *J. Am. Chem. Soc.* **113**:686 (1991); Mammen *et al.*, *J. Med. Chem.* **38**:4179 (1995)).

Moreover, polymers provide a number of advantages as the framework component, because the characteristics of the polymer can be varied, modulated and controlled as desired. For instance, characteristics which can be varied and controlled include, but are not limited to, conformal flexibility; solubility; hydrophilicity; modulation of conformation and flexibility in solution through variations in temperature and ionic strength, *etc.* As such, the use of polymers readily allows for the modulation of various physical properties of the framework. Additionally, the characteristics of the polymers can be designed to vary the flexibility of the polymer, the distance between the functional groups (*e.g.*, bioactive sidechains), the length of the spacer group or linker between the polymer backbone and the functional groups, *etc.*

The chemistry of high molecular weight polymers is a well-developed science, and organic polymers provide a very important class of compounds to use for nucleic acid conjugates. Such compounds have high molecular weights, can present very large numbers of copies of the nucleic acid and can present more than one nucleic acid simultaneously.

In preferred embodiments, modified polymeric materials for use in the present invention have low antigenicity and low toxicity. In other preferred

embodiments, polymeric frameworks are selected to be compatible with water, to have various molecular weights and to be capable of having a range of different groups attached to the polymer backbone. Polymer backbones of the present invention can also be selected for ease of synthesis.

Intrinsically biocompatible polymers containing functional groups appropriate for the addition of sidechains are preferred (Shalaby, *et al.*, *J. Polymers of Biological and Biomedical Significance (ACS Symposium Series 540)*; American Chemical Society: Washington, DC, 1994). Exemplary polymers include, but are not limited to, polyethylene oxide or polyethyleneglycol (Harris, J. M., *POLY(ETHYLENE GLYCOL) CHEMISTRY: BIOTECHNICAL AND BIOMEDICAL APPLICATIONS*; Plenum: New York, 1992; Horton, D., *ADVANCES IN CARBOHYDRATE CHEMISTRY AND BIOCHEMISTRY*; Academic Press: San Diego, 1995) as well as derivatives of acrylamide and N-vinylpyrrolidone, linked oligomers of oligoethylene glycol, linked oligomers of dextran and others.

Other preferred framework components for use in the methods of the present invention have demonstrated utility, for example, as plasma extenders, drug excipients or binders, food additives or as inert or erodible materials used *in vivo*. For example, poly(ethylene glycol), poly(lactic acid), poly(glycolic acid) and poly(vinyl pyrrolidone) can be used within the framework component in the methods of the present invention.

Preferred polymers contain reactive groups, such as amines, carboxylic acids, etc. A number of synthetic and naturally-occurring polymers containing carboxylic acid functionality (or capable of being suitably modified) have been previously used *in vivo*. Such polymers are capable of derivatization to facilitate linkage of the nucleic acid-containing subunit groups, described herein. Polymers containing internally cyclized carboxylic acid functionality, such as anhydride or succinimide groups, are also preferred. Other preferred polymers include, but are not limited to, derivatives of maleic anhydride

and malic acid. Exemplary copolymers include, but are not limited to, styrene-maleic anhydride and alpha-olefin-maleic acid copolymers (such as divinylether-maleic acid). In other embodiments, sodium carboxymethylcellulose, chondroitin sulfate and poly(methacrylate/acrylate) materials can be used. In still other embodiments, polymers without activated carboxylic acids can be used, such as dextran sulfate.

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Other exemplary polymeric framework components include, but are not limited to, poly(ester), poly(anhydride), poly(carbohydrate), polyols, poly(acrylate), poly(methacrylate), poly(ether) and poly(amino acid). Still other exemplary polymeric frameworks include, but are not limited to, polyamines, poly(glutamic acid), poly(aspartic acid), dextran, dextran sulfate, poly(maleic anhydride-co-vinyl ether), poly(succinimide), poly(acrylic anhydride), poly(ethylene glycol), poly(lactic acid), poly(glycolic acid), poly(amine), poly(vinyl pyrrolidone), poly(styrene-maleic anhydride), alpha-maleic acid, hyalouronic acid, sodium carboxymethylcellulose, chondroitin sulfate, poly(acrylate), poly(acrylamide), poly(glycerol) and starch. Table 1 sets forth a list of representative polymers useful in practicing the present invention.

**Table 1. Exemplary polymers**

Poly(ethylene glycols)	Poly(ethylene-vinyl acetate)
Poly(amides)	Poly(acrylamides)
poly(peptides)	Poly(urethanes)
Poly(aminoacids)	Poly(methacrylates)
poly(aspartic acid)	Poly(acrylates)
Poly(glutamic acid)	Poly(maleic acid)
poly(lysine), others	copolymers)
proteins(gelatins)	Poly(anhydrides)
Poly(esters)	Poly(orthoester)
poly(lactic acid),	
polylactide	
poly(glycolide)	
poly(caprolactone)	
poly(tartrate)	
Polysaccharides	
cellulose	
alginates	
starch	
Dextran derivatives	
Poly(N-vinylpyrrolidone)	

It will be appreciated by the skilled artisan that substantially any polymeric material which is capable of presenting a plurality of functional groups is suitable for use in the present invention. Polymers can be modified, *e.g.*, as described above or by derivatization, *e.g.*, with bifunctional cross-linking reagents, to provide functionalities suitable for presenting functional groups as described in more detail below.

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#### *Noncovalent Framework Components*

A plurality of ethylene-containing nucleic acids can also be joined to a non-covalent framework. Exemplary noncovalent frameworks include, but are not limited to, liposomes, micelles, colloids, protein aggregates, modified cells, and modified viral particles. For example, functional and/or ancillary groups can be tethered to the head groups of molecules in liposomes, membranes or surfaces (*see, e.g.*, Kingery-Wood

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*et al.*, *J. Am. Chem. Soc.* 114:7303-7305 (1992); Spevak *et al.*, *J. Am. Chem. Soc.* 115:1146-1147 (1993); Spevak *et al.*, *J. Med Chem.* 39:1018-1020 (1996)).

Liposomes and micelles are art-recognized terms and include macroscopic particles made up of aggregates of surfactants. In one embodiment, the compound of the invention can present groups on a liposome or micelle (Spevak *et al.*, *J. Am. Chem. Soc.* 115:1146-1147 (1993); Charych *et al.*, *Chem. & Biol.* 3:113-120 (1996)). This system closely mimics the shape of the target cell, and can be designed to present a surface that closely matches that of the target cell both in terms of group type and group density. For example, lipid molecules containing functional groups (*e.g.*, neuraminic acid (NeuAc)) as polar head groups can be reconstituted into liposomes. Liposomes have favorable biocompatibility and are fairly easy to synthesize.

In other embodiments, biological particles including, for example, modified cells or modified viruses can be used as the framework component for presentation of the functional group. Thus, proteins, peptides, polysaccharides, fragments of cell membranes, or modified intact cells (*e.g.*, erythrocytes), modified bacterial cells or modified viruses can be used as the framework component in certain embodiments.

#### *Activated Framework Component*

As used herein, the term "activated framework component" refers to a framework component, as described above, containing groups that can be activated, by means of an "activating group," and subsequently reacted with at least one functional group, ancillary group and/or spacer group that includes a group that is reactive with the ethylene-derivatized nucleic acid. Appropriate functionality includes, for example, carboxyl (acid form and salts), hydroxyl, sulfhydryl, amide, carbamate, amino, ketone, aldehyde, olefin, aromatic, *etc.* The polymers can be activated prior to exposing them to the functional group ("preactivation"), or can be activated in the presence of the functional group (*"in situ"*).

The activation step can entail derivatizing the polymer with groups capable of undergoing reactions with nucleophiles or electrophiles (*e.g.*, forming active esters, halo derivatives, *etc.*). Further, it is within the scope of the present invention to activate polymers such that they are able to participate in dipolar additions (*e.g.*, 1,3- and 1,4-dipolar addition), cycloaddition reactions (*e.g.*, Diels-Alder type reactions) and polymerization reactions by cationic, anionic or radical initiated mechanisms.

Carboxyl groups can be activated for reaction with nucleophiles by the use of, for example, cyclic or linear anhydrides, activated esters (e.g., N-hydroxysuccinimide, nitrophenol, 4-hydroxy-3-nitrobenzene sulfonic acid, *etc.*), acid chlorides, imidazolides (e.g., from carbonyldiimidazole), carboxylic acid and esters. Carboxylic acid containing polymers may also be activated by forming adducts between the carboxyl group and agents such as, dicyclohexylcarbodiimide, 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide, alkyl chloroformates, chlorosilanes, pyridinium salts  $\text{Bu}_3\text{N}$ , *etc.*

The selection of appropriate activating groups for the carboxyl functionality will be apparent to those of skill in the art. It will be similarly apparent to those of skill in the art which reaction systems will be amenable to, or will require, *in situ* activation or preactivation.

Hydroxyl groups can be activated by the use of carbonates formed by reaction with, for example, alkyl or aryl haloformates (e.g., vinylchloroformate, i-butylchloroformate, p-nitrophenylchloroformate, *etc.*), cyanogen bromide or phosgene. In aspects utilizing polymers containing vic-diol groups (e.g., dextran and other polysaccharides) oxidation using periodate compounds can be used to provide reactive carbonyl moieties on the polymeric backbone. The carbonyl moieties are then reacted with activating groups, such as vinylamine derivatives (e.g., 4-aminostyrene, *etc.*).

In a preferred embodiment, a hydroxyl-containing polymer is reacted with an active ester of acrylic acid, such as the N-hydroxysuccinimide ester to form a cleavable ester linkage between the polymer framework and the nucleic acid. (In another embodiment, the hydroxyl groups are reacted with an agent, such as vinyl bromide, vinyl benzyl chloride, *etc.*). Additional methods of activating polymers bearing hydroxyl groups will be apparent to those of skill in the art.

Polymers bearing sulphhydryl groups can be activated using dithiobispyridyl compounds such as, for example, 2,2'-dithiobis(5-nitropyridine), 2,2'-dithiobis(pyridine), *etc.* Additional methods of use in activating sulphhydryl-bearing polymers will be apparent to those of skill in the art.

It will be appreciated by those of skill in the art that the above activation reactions are set forth as examples only and that many further alternatives to these schemes exist.

In preferred embodiments, the polymeric framework is polycationic. Polycationic groups have been shown to facilitate the entry of polyanionic nucleic acids into cell. Useful polycationic motifs include polyamines such as polylysine.

Additional polymers, linker groups functional groups and combinations thereof within both the scope and spirit of the present invention will be apparent to those of skill in the art.

5                   (ii). Nucleic acids

Any nucleic acid can be used to construct the compositions of the invention. Particularly preferred naturally occurring nucleic acid molecules include genomic deoxyribonucleic acid (DNA) and genomic ribonucleic acid (RNA), as well as the several different forms of the latter, *e.g.*, messenger RNA (mRNA), transfer RNA (tRNA), and ribosomal RNA (rRNA). Also included are the different DNAs which are complementary (cDNA) to the different RNAs. Synthetic DNA or a hybrid thereof with naturally occurring DNA, is also encompassed within the scope of the instant disclosure.

10                  The nucleic acid compositions used in the present invention may be either single-stranded or double-stranded, may be linear or circular, *e.g.*, a plasmid, and are either oligo- or poly-nucleotides. They may comprise a single base or base pair, or may include as many as 20 thousand bases or base pairs (20 kb), or more.

15                  In addition to these naturally occurring materials, the nucleic acid compositions used in the present invention can also include synthetic compositions, *e.g.*, nucleic acid analogs, synthetic nucleic acids. These have been found to be particularly useful in antisense methodology, which is the complementary hybridization of relatively short oligonucleotides to single-stranded RNA or single-stranded DNA, such that the normal, essential functions of these intracellular nucleic acids are disrupted. *See, e.g.*, Cohen, OLIGONUCLEOTIDES: ANTISENSE INHIBITORS OF GENE EXPRESSION, CRC Press, Inc., Boca Raton, Fla. (1989).

20                  The nucleic acid composition to be transferred to a target cell in accordance with the present invention preferably has an appropriate open reading frame and promoter to express a protein, as well as any other regulatory sequences which may be appropriate to expression. Nucleic acid compositions to be delivered by means of the methods of the present invention can be designed and constructed so as to be appropriate for the particular application desired, all of which is well within the ordinary skill of the artisan in this field.

25                  The nucleic acid molecules which are delivered to cells using the multifunctional molecular complex and methods of the present invention may, for example, serve as: 1) genetic templates for proteins that function as prophylactic and/or

therapeutic immunizing agents; 2) replacement copies of defective, missing or non-functioning genes; 3) genetic templates for therapeutic proteins; 4) genetic templates for antisense molecules and as antisense molecules *per se*; or 5) genetic templates for ribozymes.

5 In the case of nucleic acid molecules which encode proteins, the nucleic acid molecules preferably comprise the necessary regulatory sequences for transcription and translation in the target cells of the individual animal to which they are delivered.

10 In the case of nucleic acid molecules which serve as templates for antisense molecules and ribozymes, such nucleic acid molecules are preferably linked to regulatory elements necessary for production of sufficient copies of the antisense and ribozyme molecules encoded thereby respectively.

The nucleic acid molecules are free from retroviral particles and are preferably provided as DNA in the form of plasmids.

15 In some cases, it may be desirable to use constructs that produce long term transgene effects *in vivo*, either by integration of the transgene into host cell genomic DNA at high levels or by persistence of the transgene in the nucleus of cells *in vivo* in stable, episomal form. Integration of the transgene into genomic DNA of host cells *in vivo* may be facilitated by administering the transgene in a linearized form (either the coding region alone, or the coding region together with 5' and 3' regulatory sequences, but without any plasmid sequences present). It is possible to further increase the incidence of transgene integration into genomic DNA by incorporating a purified retroviral enzyme, such as the HIV-1 integrase enzyme, into the lipid carrier-DNA complex. Appropriate flanking sequences are placed at the 5' and 3' ends of the transgene DNA. These flanking sequences have been shown to mediate integration of the HIV-1 DNA into host cell  
20 genomic DNA in the presence of HIV-1 integrase. Alternatively, duration of transgene expression *in vivo* can be prolonged by the use of constructs that contain non-transforming sequences of a virus such as Epstein-Barr virus, and sequences such as oriP and EBNA-1, which appear to be sufficient to allow heterologous DNA to be replicated as a plasmid in mammalian cells (Buhans *et al.*, *Cell* 62:955 (1986)).

25 30 The nucleic acid constructs for use in the invention include several forms, depending upon the intended use of the construct. Thus, the constructs include, for example, vectors, transcriptional cassettes, expression cassettes and plasmids. The transcriptional and translational initiation region (also sometimes referred to as a "promoter,"), preferably comprises a transcriptional initiation regulatory region and a

translational initiation regulatory region of untranslated 5' sequences, "ribosome binding sites," responsible for binding mRNA to ribosomes and translational initiation. It is preferred that all of the transcriptional and translational functional elements of the initiation control region are derived from or obtainable from the same gene. In some 5 embodiments, the promoter will be modified by the addition of sequences, such as enhancers, or deletions of nonessential and/or undesired sequences. "Obtainable," as used herein, refers to a promoter having a DNA sequence sufficiently similar to that of a native promoter to provide for the desired specificity of transcription of a DNA sequence of interest. It includes natural and synthetic sequences as well as sequences which may be a 10 combination of synthetic and natural sequences.

For the transcriptional initiation region, or promoter element, any region 15 may be used with the proviso that it provides the desired level of transcription of the DNA sequence of interest. The transcriptional initiation region may be native to or homologous to the host cell, and/or to the DNA sequence to be transcribed, or foreign or heterologous to the host cell and/or the DNA sequence to be transcribed. "Foreign to the host cell," as used herein, refers to sequences in which the transcriptional initiation region is not found in the host into which the construct comprising the transcriptional initiation region is to be inserted. "Foreign to the DNA sequence," as used herein, refers to a 20 sequence in which a transcriptional initiation region that is not normally associated with the DNA sequence of interest. Efficient promoter elements for transcription initiation include, for example, the SV40 (simian virus 40) early promoter, the RSV (Rous sarcoma virus) promoter, the Adenovirus major late promoter, and the human CMV (cytomegalovirus) immediate early 1 promoter.

Inducible promoters also find use with the subject invention where it is 25 desired to control the timing of transcription. Examples of promoters include but are not limited to those obtained from a beta-interferon gene, a heat shock gene, a metallothionein gene or those obtained from steroid hormone-responsive genes, including insect genes such as that encoding the ecdysone receptor. Such inducible promoters can be used to regulate transcription of the transgene by the use of external stimuli such as 30 interferon or glucocorticoids. Since the arrangement of eukaryotic promoter elements is highly flexible, combinations of constitutive and inducible elements also can be used. Tandem arrays of two or more inducible promoter elements may increase the level of induction above baseline levels of transcription which can be achieved when compared to

the level of induction above baseline which can be achieved with a single inducible element.

Generally, the regulatory sequence comprises DNA up to about 1.5 Kb 5' of the transcriptional start of a gene, but this sequence can be significantly smaller. The 5 regulatory sequence can be modified at a position corresponding to the first codon of the desired protein by site-directed mutagenesis (Kunkel, *Proc. Natl. Acad. Sci. (USA)* 82:488-492 (1985)) or by introduction of a convenient linker oligonucleotide by ligation, if a suitable restriction site is found near the N-terminal codon. In a preferred embodiment, a coding sequence with a compatible restriction site may be ligated at the 10 position corresponding to codon #1 of the gene. This substitution may be inserted in such a way that it completely replaces the native coding sequence and thus the substituted sequence is flanked at its 3' end by the gene terminator and polyadenylation signal.

Transcriptional enhancer elements optionally are included in the expression cassette. "Transcriptional enhancer elements," as used herein, refer to DNA sequences which are primary regulators of transcriptional activity and which can act to increase transcription from a promoter element. These elements generally do not have to be in the 5' orientation with respect to the promoter in order to enhance transcriptional activity. The combination of promoter and enhancer element(s) used in a particular expression cassette can be selected by one skilled in the art to maximize specific effects. 20 Different enhancer elements can be used to produce a desired level of transgene expression in a wide variety of tissue and cell types. For example, the human CMV immediate early promoter-enhancer element can be used to produce high-level transgene expression in many different tissues *in vivo*.

Examples of other enhancer elements which confer a high level of 25 transcription on linked genes in a number of different cell types from many species include enhancers from SV40 and RSV-LTR. The SV40 and RSV-LTR are essentially constitutive. They may be combined with other enhancers which have specific effects, or the specific enhancers may be used alone. Thus, where specific control of transcription is desired, efficient enhancer elements that are active only in a tissue-, developmental-, or 30 cell-specific fashion include immunoglobulin, interleukin-2 (IL-2) and beta -globin enhancers are of interest. Tissue-, developmental-, or cell-specific enhancers can be used to obtain transgene expression in particular cell types, such as B-lymphocytes and T-lymphocytes, as well as myeloid, or erythroid progenitor cells. Alternatively, a tissue-specific promoter such as that derived from the human cystic fibrosis transmembrane

conductance regulator (CFTR) gene can be fused to a very active, heterologous enhancer element, such as the SV40 enhancer, in order to confer both a high level of transcription and tissue-specific transgene transcription. In addition, the use of tissue-specific promoters, such as LCK, may allow targeting of transgene transcription to T lymphocytes. Tissue specific transcription of the transgene may be important, particularly in cases where the results of transcription of the transgene in tissues other than the target tissue would be deleterious.

Tandem repeats of two or more enhancer elements or combinations of enhancer elements may significantly increase transgene expression when compared to the use of a single copy of an enhancer element; hence enhancer elements find use in the expression cassette. The use of two different enhancer elements from the same or different sources flanking or within a single promoter can in some cases produce transgene expression in each tissue in which each individual enhancer acting alone would have an effect, thereby increasing the number of tissues in which transcription is obtained. In other cases, the presence of two different enhancer elements results in silencing of the enhancer effects. Evaluation of particular combinations of enhancer elements for a particular desired effect or tissue of expression is within the level of skill in the art.

Although generally it is not necessary to include an intron in the expression cassette, an intron comprising a 5' splice site (donor site) and a 3' splice site (acceptor site) separated by a sufficient intervening sequence to produce high level, extended *in vivo* expression of a transgene administered iv or ip can optionally be included. Generally, an intervening sequence of about 100 bp produces the desired expression pattern and/or level, but the size of the sequence can be varied as needed to achieve a desired result. The optional intron placed 5' to the coding sequence results in high level extended *in vivo* expression of a transgene administered iv or ip but generally is not necessary to obtain expression. Optimally, the 5' intron specifically lacks cryptic splice sites which result in aberrantly spliced mRNA sequences. If used, the intron splice donor and splice acceptor sites, arranged from 5' to 3' respectively, are placed between the transcription initiation site and the translational start codon.

Alternatively, the intervening sequence may be placed 3' to the translational stop codon and the transcriptional terminator or inside the coding region. The intron can be a hybrid intron with an intervening sequence or an intron taken from a genomic coding sequence. An intron 3' to the coding region, particularly one of less than

100 bp, or any intron which contains cryptic splice sites may under certain condition substantially reduce the level of transgene expression produced *in vivo*. A high level of *in vivo* expression of a transgene can also be achieved using a vector that lacks an intron. Such vectors therefore are of particular interest for *in vivo* transfection.

5 Downstream from and under control of the transcriptional initiation regulatory regions is, preferably, a multiple cloning site for insertion of a nucleic acids sequence of interest which will provide for one or more alterations of host genotype and modulation of host phenotype. Conveniently, the multiple cloning site may be employed for a variety of nucleic acid sequences in an efficient manner. The nucleic acid sequence 10 inserted in the cloning site may have any open reading frame encoding a polypeptide of interest, for example, an enzyme, with the proviso that where the coding sequence encodes a polypeptide of interest, it should lack cryptic splice sites which can block production of appropriate mRNA molecules and/or produce aberrantly spliced or abnormal mRNA molecules. The nucleic acid sequence may be DNA; it also may be a sequence complementary to a genomic sequence, where the genomic sequence may be one or more of an open reading frame, an intron, a non-coding leader sequence, or any other sequence where the complementary sequence will inhibit transcription, messenger RNA processing, for example splicing, or translation.

15 A number of nucleic acid sequences are of interest for use in *in vivo* gene therapy. Where the nucleic acid codes for a polypeptide, the polypeptide may be one 20 which is active intracellularly, a transmembrane protein, or it may be a secreted protein. It may also code for a mutant protein, for example, which is normally secreted, but which has been altered act intracellularly. The nucleic acid may also be a DNA sequences 25 coding for mRNA (antisense or ribozyme sequences such as those to HIV-REV or BCR-ABL sequences) or for proteins such as transdominant negative mutants, which specifically prevent the integration of HIV genes into the host cell genomic DNA, replication of HIV sequences, translation of HIV proteins, processing of HIV mRNA or virus packaging in human cells; the LDL (low density lipoprotein) receptor, which 30 specifically lowers serum cholesterol; and proteins such as granulocyte macrophage colony stimulating factor (GM-CSF) which can stimulate the production of white blood cells from the bone marrow of immunocompromised patients and produce significant anti-tumor activity or cystic fibrosis transmembrane conductance regulator (CFTR) for the treatment of cystic fibrosis. These and other beneficial (therapeutic) nucleic acid sequences can be expressed in appropriate cells *in vivo* using this invention.

Examples of beneficial therapeutic nucleic acid sequences are those encoding molecules having superoxide dismutase activity or catalase activity to protect the lung from oxidant injury; endothelial prostaglandin synthase to produce prostacyclin and prostaglandin E2; and antiprotease alpha-1 antitrypsin. Thus, this approach could 5 dramatically improve the treatment of acquired immune deficiency syndrome (AIDS), cystic fibrosis, cancer, heart disease, autoimmune diseases and a variety of life threatening infections. For the treatment of AIDS, anti-TAT, REV, TAR or other critical anti-HIV sequences can be used, particularly for expression of the appropriate coding sequences in T lymphocytes, macrophages and monocytes which can be achieved 10 following iv administration of the appropriate coding sequences; expression of wild-type CFTR gene in the lungs of cystic fibrosis patients (*see, Collins, Science 256:774-783 (1992)*); antisense sequences to over-expressed, transforming oncogenes, such as myc or ras in tumors; and genes which block activity of activated T cell clones which attack 15 myelin in multiple sclerosis or other targets in autoimmune diseases. A T-cell lymphocyte clone activated to recognize and attack myelin can be targeted by using an anti-sense sequence, ribozyme sequence or transgene coding for a transdominant negative mutant which specifically blocks surface expression on the T-cell of T-cell receptor components which mediate recognition and/or attack of myelin-sheathed cells.

The choice of termination region employed will primarily be one of convenience, since termination regions appear to be relatively interchangeable. The 20 termination region may be native to the intended nucleic acid sequence of interest, or it may be derived from another source. Convenient termination regions are available and include the 3' end of a gene terminator and polyadenylation signal from the same gene from which the 5' regulatory region is obtained. Adenylation residues, preferably more 25 than 32 and up to 200 or more as necessary may be included in order to stabilize the mRNA. Alternatively, a terminator and polyadenylation signal from different gene/genes may be employed with similar results. Specific sequences which regulate post-transcriptional mRNA stability may optionally be included. For example, certain polyA sequences (Volloch *et al., Cell 23:509 (1981)*) and beta -globin mRNA elements can 30 increase mRNA stability, whereas certain AU-rich sequences in mRNA can decrease mRNA stability (Shyu *et al., Genes and Devel. 3:60 (1989)*). In addition, AU regions in 3' non-coding regions may be used to destabilize mRNA if a short half-life mRNA is desirable for the gene of interest.

The construct may additionally include sequences for selection, such as a neomycin resistance gene or a dihydrofolate reductase gene and/or signal sequences to regenerate recombinant proteins that are targeted to different cellular compartments or secreted when the wild type sequence is not. Any of a variety of signal sequences may be used which are well-known to those skilled in the art. These signal sequences may allow generation of new vaccine strategies or produce soluble antagonists directed against specific cell surface receptors such as transformed oncogenes. The sequences for selection may be on a separate plasmid and cotransfected with the plasmid carrying the therapeutic nucleic acid. Where a carrier is used, the selection plasmid may be complexed to a different carrier or to the same carrier as the therapeutic plasmid.

The recombinant coding-sequence flanked at its 5' end by the promoter and regulatory sequences and at its 3' end by a terminator and regulatory sequences may be introduced into a suitable cloning plasmid (e.g., pUC18, pSP72) for use in direct DNA uptake in host cells following introduction into the host. The nucleic acid construct also may be complexed with a carrier such as lipid carriers, particularly cationic lipid carriers.

As discussed above, the nucleic acids can be either single- or double-stranded or combinations thereof. In a preferred embodiment the first nucleic acid tethered to the polymer is hybridized to a second nucleic acid. In a further preferred embodiment, the first nucleic acid is a single-stranded nucleic acid. In another preferred embodiment, the first nucleic acid is a double-stranded nucleic acid. In these embodiments, the second nucleic acid can be either single- or double stranded, forming either duplexes or triplexes with the immobilized first nucleic acid, respectively.

#### (b). Particles

In another aspect, the invention provides a polymeric particle including a first subunit. The first subunit includes a first nucleic acid. The first subunit is incorporated into the polymer using a first subunit precursor. The first subunit precursor includes an ethylene-containing moiety. The embodiments of this aspect are substantially similar to those discussed above in the context of the nucleic acids and substantially water-soluble polymers and many of the polymers discussed above can be used to form the particles of the invention.

Many different types of microparticles for drug delivery are known in the art. Of particular relevance to the present invention are those microparticles that are prepared from ethylene-containing monomers, such as acrylate derivatives.

Microparticles prepared from acrylic monomers have attracted a great deal of attention as colloidal drug carriers because of their ease of preparation. Cyanoacrylate particles are both lysosometric, biodegradable and biocompatible ( Couvreur *et al.*, *J. Pharmacol. Sci.* **68**:1521 (1979)). Cyanoacrylate particles have been used in a number of different applications, such as ocular drug delivery, carriers of monoclonal antibodies and for targeting chemotherapeutic agents to cancer cells (Tuncel *et al.*, *J. Biomed. Mat. Res.* **29**:721 (1995)).

Many methods have been developed to form discrete nanoparticles from acrylate monomers. These include, for example, aqueous anionic polymerization at a low pH (see, for example, Kreuter *et al.*, *Int. J. Pharmacol.* **16**:105 (1983); Couvreur *et al.*, *J. Pharm. Pharmacol.* **31**:331 (1979); Douglas *et al.*, *J. Colloid. Interface Sci.* **101**:149 (1984)). Cyanoacrylate microspheres have also been prepared by dispersion polymerization in an acidic aqueous solution (Tuncel, *supra*; Sjöholm *et al.*, *J. Pharm. Exp. Ther.* **211**:656 (1979)). Methods have also been developed to form a population of microspheres, the members of which are of practically a single size. For example, Amsden (*Pharm. Res.* **16**:140 (1999)) has developed a method involving the injection of a solution to be emulsified and formed into microspheres into a stabilizing solution flowing past an injection point. Thus, using methods such as that disclosed by Amsden microspheres having a very narrow diameter distribution are produced.

In an exemplary embodiment, the nucleic acid-containing subunit includes an acrylamide moiety. The nucleic acid-acrylamide conjugate is combined with bisacrylamide (3:1 w/w) in an aqueous buffer (e.g., sodium phosphate, pH 7.4). The solution is optionally deoxygenated by bubbling nitrogen gas through the solution. A catalyst, such as ammonium peroxodisulfate in water is added and the aqueous component is homogenized in an organic medium comprising a mixture of toluene and chloroform (4:1) containing a detergent, such as Pluronic F-68. Polymerization of the resulting emulsion is initiated by the addition of N,N',N'',N'''-tetramethylethylenediamine. The resulting suspension is stirred for approximately one-half hour and the phases are separated by centrifugation. The organic phase is removed and the microparticles located at the bottom of the aqueous phase are repeatedly washed with an aqueous buffer.

The resulting microparticles are characterized by methods well known in the art. Using standard techniques, the half-life of the particle in the blood can be determined and the distribution of the particles in the body is determined by, for example,

injecting radioactively labeled microspheres and counting the individual organs after sacrifice of the experimental animal (Sjöholm *et al.*, *supra*). The particle size is also determined by art recognized methods, such as scanning electron microscopy (Höglund *et al.*, *J. Gen. Virol.* 21:359 (1973)). Other methods of preparing and characterizing microparticles according to the present invention will be apparent to those of skill in the art.

The particles of the invention can be of substantially any useful size for a particular application. In a preferred embodiment, the particles range from about 0.001  $\mu\text{m}$  to about 100  $\mu\text{m}$ , more preferably from about .1  $\mu\text{m}$  to about 10  $\mu\text{m}$  in diameter.

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**(c). Tissue targeted agents and promoters of cellular uptake**

In the methods of the invention, the amount of transfection desired is that which will result in a therapeutic effect, *i.e.* prevention, palliation, and/or cure of an animal or human disease ("*in vivo*" gene therapy). Optionally, the carrier molecule and/or construct of the invention may provide for targeting and/or expression in a particular cell type or types.

The size, nature and specific sequence of the nucleic acid composition to be transferred to the target cell can be optimized for the particular application for which it is intended, and such optimization is well within the skill of the artisan in this field. However, the nature of the target cells within the individual into which it is desired to transfer a nucleic acid composition, may have a significant bearing on the choice of the particular multifunctional molecular complex of the present invention. For example, where it is desired to transfer nucleic acid molecules to target cells by injecting them intramuscularly to evoke an immune response, it will be found that this transfer can be effected by use of a multifunctional molecular complex of the present invention, as defined above, comprising a cationic polyamine to which is attached, as the endosome membrane disruption promoting component, a lipophilic long chain alkyl group as defined above.

Another approach in the art to delivery of genetic material to target cells is one that takes advantage of natural receptor-mediated endocytosis pathways that exist in such cells. Several cellular receptors have been identified heretofore as desirable agents by means of which it is possible to achieve the specific targeting of drugs, and especially macromolecules and molecular conjugates serving as carriers of genetic material of the

type with which the present invention is concerned. These cellular receptors allow for specific targeting by virtue of being localized to a particular tissue or by having an enhanced avidity for or activity in a particular tissue. See, e.g., J. L. Bodmer *et al.*, *Meth. Enzymol.* **112**:298-306 (1985). This affords the advantages of lower doses or 5 significantly fewer undesirable side effects.

One of the better known examples of a cell and tissue selective receptor is the asialoglycoprotein receptor present in hepatocytes. The asialoglycoprotein receptor is an extracellular receptor with a high affinity for galactose, especially tri-antennary oligosaccharides, i.e., those with three somewhat extended chains or spacer arms having 10 terminal galactose residues; see, e.g., Lodish, *TIBS* **16**:374-77 (1991). This high affinity receptor is localized to hepatocytes and is not present in Kupffer cells; allowing for a high degree of selectivity in delivery to the liver.

It has also been proposed in the art of receptor-mediated gene transfer that in order for the process to be efficient *in vivo*, the assembly of the DNA complex should result in condensation of the DNA to a size suitable for uptake via an endocytic pathway. 15 See, e.g., Perales *et al.*, *Proc. Nat. Acad. Sci. USA* **91**:4086-4090 (1994).

An alternative method of providing cell-selective binding is to attach an entity with an ability to bind to the cell type of interest; commonly used in this respect are antibodies which can bind to specific proteins present in the cellular membranes or outer regions of the target cells. Alternative receptors have also been recognized as useful in facilitating the transport of macromolecules, such as biotin and folate receptors; see, Low 20 *et al.*, WO 90/12095, published 18 Oct. 1990; Low *et al.*, WO 90/12096, published 18 Oct. 1990; Low *et al.*, U.S. Pat. No. 5,108,921, Apr. 28, 1992; Leamon *et al.*, *Proc. Nat. Acad. Sci. USA* **88**:5572-5576 (1991); transferrin receptors; insulin receptors; and 25 mannose receptors (see further below). The enumerated receptors are merely representative, and other examples will readily come to the mind of the artisan.

The conjugation of different functionalities on the same molecule has also been utilized in the art. For example, in 1988, Wu *et al.*, *J. Biol. Chem.* **263**:14621-14624 (1988) described a method for cellular receptor mediated delivery of DNA to hepatocytes. 30 This method was further described in Wu *et al.*, *Biochem.* **27**:887-892 (1988); Wu *et al.*, U.S. Pat. No. 5,166,320, Nov. 24, 1992; and Wu *et al.*, WO 92/06180, published 16 Apr. 1992. The method consists of attaching a glycoprotein, asialoorosomucoid, to poly-lysines to provide a hepatocyte selective DNA carrier. The function of the poly-lysine is to bind to the DNA through ionic interactions between the positively charged (cationic)

epsilon amino groups of the lysines and the negatively charged (anionic) phosphate groups of the DNA. Orosomucoid is a glycoprotein, which is normally present in human serum. Removal of the terminal sialic acid (N-acetyl neuraminic acid) from the branched oligosaccharides exposes terminal galactose oligosaccharides, for which hepatocyte receptors have a high affinity, as already described.

After binding to the asialoglycoprotein receptor on hepatocytes, the protein is taken into the cell by endocytosis into a pre-lysosomal endosome. The DNA, ionically bound to the poly-lysine-asialoorosomucoid carrier, is also taken into the endosome. Additional work using this delivery system, *e.g.*, that done by Wilson *et al.*, *J. Biol. Chem.* 267:11483-11489 (1992), has found that partial hepatectomy improves the persistence of the expression of the DNA delivered into the hepatocytes. The transfer of the DNA into cells by this mechanism is also significantly enhanced by the addition of cationic lipids; *see, e.g.*, Mack *et al.*, *Am. J. Med. Sci.* 307:138-143 (1994).

The use of a specific asialoglycoprotein is not required to achieve binding to the asialoglycoprotein receptor; this binding can also be accomplished with high affinity by the use of small, synthetic molecules having a similar configuration. The carbohydrate portion can be removed from an appropriate glycoprotein and be conjugated to other macromolecules; *see, e.g.*, Wood *et al.*, *Bioconj. Chem.* 3:391-396 (1992). By this procedure the cellular receptor-binding portion of the glycoprotein is removed, and the specific portion required for selective cellular binding can be transferred to another molecule.

There is an ample literature on the preparation of synthetic glycosides which can be attached to macromolecules and confer on them the ability to bind to the corresponding galactose specific receptor. The importance of branched glycosides was recognized early; *see*, Lee, *Carb. Res.* 67:509-514 (1978). Further work delineated that sugar density (Kawaguchi *et al.*, *J. Biol. Chem.* 256:2230-2234 (1981)) and spacial relationships (Lee *et al.*, *J. Biol. Chem.* 258:199-202 (1983)) are important determinants of binding potency. Reductive amination of a peptide with a branched tri-lysine amino terminus gives a ligand ending with four galactosyl residues that can be readily coupled to poly-lysine or other macromolecules and has been used to prepare DNA constructs; *see*, Plank *et al.*, *Bioconj. Chem.* 3:533-539 (1992);

Thiopropionate and thiohexanoate glycosidic derivatives of galactose have been prepared and linked to L-lysyl-L-lysine to form a synthetic tri-antennary galactose derivative. A bisacridine spermidine derivative containing this synthetic tri-antennary

galactose has been used to target DNA to hepatocytes; *see, Szoka et al.*, WO 93/19768, published 14 Oct. 1993; and Haensler *et al.*, *Bioconj. Chem.* 4:85-93 (1993).

Other means of providing cellular receptor based facilitation of gene transfer into cells using carriers have been described in the art. Antibodies specific for 5 cell surface thrombomodulin have been used with poly-lysine as a delivery system for DNA *in vitro* and *in vivo*; *see Trubetskoy et al.*, *Bioconj. Chem.* 3:323-327 (1992). The transferrin receptor has also been used to target DNA to erythroblasts, K562 macrophages and ML-60 leukemic cells; *see, Wagner et al.*, *Proc. Nat. Acad. Sci. USA* 87:3410-3414 (1990); Zenke *et al.*, *Proc. Nat. Acad. Sci. USA* 87:3655-3659 (1990); and Citro *et al.*, 10 *Proc. Nat. Acad. Sci. USA* 89:7031-7035 (1990). These studies used both small oligodeoxynucleotides as well as large plasmids.

The ability of poly-lysine to facilitate DNA entry into cells is significantly enhanced if the poly-lysine is chemically modified with hydrophobic appendages; *see Zhou et al.*, *Biochim. Biophys. Acta* 1189:195-203 (1994); complexed with cationic 15 lipids; *see Mack et al.*, *Am. J. Med. Sci.* 307:138-143 (1994) or associated with viruses. Many viruses infect specific cells by receptor-mediated binding and insertion of the viral DNA/RNA into the cell and thus this action of the virus is similar to the facilitated entry of DNA described above.

Replication-incompetent adenovirus has been used to enhance the entry of 20 transferrin-poly-lysine complexed DNA into cells; *see, Curiel et al.*, *Proc. Nat. Acad. Sci. USA* 88:8850-8854 (1991); Wagner *et al.*, *Proc. Nat. Acad. Sci. USA* 89:6099-6103 (1992); Cotton *et al.*, *Proc. Nat. Acad. Sci. USA* 89:6094-6098 (1992); and Gao *et al.*, *Hum. Gene Ther.* 4:17-24 (1993). The adenovirus enhances the entry of the poly-lysine-transferrin-DNA complex when covalently attached to the poly-lysine and when attached 25 through an antibody-binding site. There does not need to be a direct attachment of the adenovirus to the poly-lysine-transferrin-DNA complex, and it can facilitate the entry of the complex when present as a simple mixture. The poly-lysine transferrin-DNA complex provides receptor specific binding to the cells and is internalized into endosomes along with the DNA. Once inside the endosomes, the adenovirus facilitates entry of the 30 DNA/transferrin-poly-lysine complex into the cell by disruption of the endosomal compartment with subsequent release of the DNA into the cytoplasm. Replication-incompetent adenovirus has also been used to enhance the entry of uncomplexed DNA plasmids into cells without the benefit of the cell receptor selectivity conferred by the

poly-lysine-transferrin complex; *see*, Yoshimura *et al.*, *J. Biol. Chem.* **268**:2300-2303 (1993).

Synthetic peptides such as the N-terminus region of the influenza hemagglutinin protein are known to destabilize membranes and are known as fusogenic peptides. Conjugates containing the influenza fusogenic peptide coupled to poly-lysine together with a peptide having a branched tri-lysine amino terminus ligand ending with four galactosyl residues have been prepared as facilitators of DNA entry into hepatocytes; *see*, Plank *et al.*, *Bioconj. Chem.* **3**:533-539 (1992). These conjugates combine the asialoglycoprotein receptor mediated binding conferred by the tetra-galactose peptide, the endosomal disrupting abilities of the influenza fusogenic peptide, and the DNA binding of the poly-lysine. These conjugates deliver DNA into the cell by a combination of receptor mediated uptake and internalization into endosomes. This internalization is followed by disruption of the endosomes by the influenza fusogenic peptide to release the DNA into the cytoplasm. In a similar fashion, the influenza fusogenic peptide can be attached to poly-lysine and mixed with the transferrin-poly-lysine complex to provide a similar DNA carrier selective for cells carrying the transferrin receptor; *see*, Wagner *et al.*, *Proc. Nat. Acad. Sci. USA* **89**:7934-7938 (1992). Synthetically designed peptides can also be used; for example the "GALA" peptides (Subbarao *et al.*, *J. Biol. Chem.* **26**:2964-2972 (1987)) have been coupled to DNA carriers and an enhanced facilitated entry into cells was observed (Haensler *et al.*, *Bioconj. Chem.* **4**:372-379 (1993)). The cationic amphipathic peptide gramicidin S can facilitate entry of DNA into cells (Legendre *et al.*, *Proc. Nat. Acad. Sci. USA* **90**:893-897 (1993)), but also requires a phospholipid to achieve significant transfer of DNA.

Positively charged liposomes have also been widely used as carriers of DNA which facilitate entry into cells; *see*, e.g., Szoka *et al.*, WO 93/19768, published 14 Oct. 1993; Debs *et al.*, WO 93/24640, published 9 Dec. 1993; Felgner *et al.*, WO 91/16024, published 31 Oct. 1991; Felgner *et al.*, *Nature* **337**:387-388 (1989); Rose *et al.*, *BioTechniques* **10**:520-525 (1991); Bennett *et al.*, *Mol. Pharm.* **41**:1023-1033 (1992); Felgner *et al.*, *J. Biol. Chem.* **269**:2550-2561 (1994); Smith *et al.*, *Biochim. Biophys. Acta* **1154**:327-340 (1993). These carrier compositions have also included pH sensitive liposomes; *see*, Chu *et al.*, *Pharm. Res.* **7**:824-854 (1990); Legendre *et al.*, *Pharm. Res.* **9**:1253-1242 (1992).

A poly-cationic lipid has been prepared by coupling dioctadecylamidoglycine and dipalmitoyl phosphatidylethanolamine to a 5-

carboxyspermine; *see*, Behr *et al.*, *Proc. Nat. Acad. Sci. USA* **86**:6982-6986 (1989); Barthel *et al.*, *DNA and Cell Biol.* **12**:553-560 (1993); Loeffler *et al.*, *Meth. Enzymol.* **217**:599-618 (1993); Behr *et al.*, U.S. Pat. No. 5,171,678, Dec. 15, 1992. These lipophilic-spermines are very active in transferring DNA through cellular membranes.

5 Combinations of lipids have been used to facilitate the transfer of nucleic acids into cells. For example, in U.S. Pat. No. 5,283,185 there is disclosed such a method, which utilizes a mixed lipid dispersion of a cationic lipid with a co-lipid in a suitable solvent. The lipid has a structure, which includes a lipophilic group derived from cholesterol, a linker bond, a linear alkyl spacer arm, and a cationic amino group; and the 10 co-lipid is phosphatidylcholine or phosphatidylethanolamine.

Macrophages have receptors for both mannose and mannose-6-phosphate, which can bind to and internalize molecules displaying these sugars. The molecules are internalized by endocytosis into a pre-lysosomal endosome. This internalization has been used to enhance entry of oligonucleotides into macrophages using bovine serum albumin modified with mannose-6-phosphate and linked to an oligodeoxynucleotide by a disulfide bridge to a modified 3' end; *see*, Bonfils *et al.*, *Nucl. Acids Res.* **20**:4621-4629 (1992). Similarly, oligodeoxynucleotides modified at the 3' end with biotin were combined with mannose-modified streptavidin, and were also found to have facilitated entry into macrophages; *see*, Bonfils *et al.*, *Bioconj. Chem.* **3**:277-284 (1992).

20 Various peptides and proteins, many of which are naturally occurring, have been shown to have receptors on cell surfaces that once they are attached thereto, allow them to become internalized by endocytosis. Materials bound to these receptors are delivered to endosomal compartments inside the cell. Examples include insulin, vasopressin, low-density lipoprotein, epidermal growth factor and others. This 25 internalization has also been used to facilitate entry of DNA into cells; *e.g.*, insulin has been conjugated to polylysine to provide facilitated DNA entry into cells possessing an insulin receptor; *see*, Huckett *et al.*, *Biochem. Pharmacol.* **40**:253-263 (1990).

In another preferred embodiment, the present invention provides 30 compositions that include a moiety that enhances cellular uptake. Many of the moieties discussed above not only aid in targeting the nucleic acids to a particular cell, but also enhance the uptake of the constructs by the cell. Moreover, delivery vehicles in which the composition of the invention is dissolved and/or suspended are known that enhance cellular uptake. Examples of such a delivery vehicles are provided in Edwards *et al.*, U.S. Patent No. 5,985,320, November 16, 1999.

## 2. Pharmaceutical Formulations

The invention also provides pharmaceutical compositions comprising one or more compounds of this invention in association with a pharmaceutically acceptable carrier. Preferably these compositions are in unit dosage forms such as tablets, pills, capsules, powders, granules, sterile parenteral solutions or suspensions, metered aerosol or liquid sprays, drops, ampoules, auto-injector devices or suppositories; for oral, parenteral, intranasal, sublingual or rectal administration, or for administration by inhalation or insufflation. Alternatively, the compositions may be presented in a form suitable for once-weekly or once-monthly administration; for example, an insoluble salt of the active compound, may be adapted to provide a depot preparation for intramuscular injection. An erodible polymer containing the active ingredient may be envisaged. For preparing solid compositions such as tablets, the principal active ingredient is mixed with a pharmaceutical carrier, e.g. conventional tableting ingredients such as corn starch, lactose, sucrose, sorbitol, talc, stearic acid, magnesium stearate, dicalcium phosphate or gums, and other pharmaceutical diluents, e.g. water, to form a solid preformulation composition containing a homogeneous mixture of a compound of the present invention, or a pharmaceutically acceptable salt thereof.

When referring to these preformulation compositions as homogeneous, it is meant that the compound of the invention is dispersed substantially evenly throughout the composition so that the composition may be readily subdivided into equally effective unit dosage forms such as tablets, pills and capsules.

The liquid forms in which the novel compositions of the present invention may be incorporated for administration orally or by injection include aqueous solutions, suitably flavored syrups, aqueous or oil suspensions, and flavored emulsions with edible oils such as cottonseed oil, sesame oil, coconut oil or peanut oil, as well as elixirs and similar pharmaceutical vehicles. Suitable dispersing or suspending agents for aqueous suspensions include synthetic and natural gums such as tragacanth, acacia, alginate, dextran, sodium carboxymethylcellulose, methylcellulose, polyvinyl-pyrrolidone or gelatin.

Any physiologically acceptable medium may be employed for administering the compositions of the invention, such as deionized water, saline, phosphate-buffered saline, 5% dextrose in water, and the like, depending upon the route of administration. Other components may be included in the formulation such as buffers,

stabilizers, biocides, *etc.* These components have found extensive exemplification in the literature and need not be described in particular here.

### **3. Methods**

5        The present invention further provides methods for delivering a nucleic acid composition into a cell. The method comprises the step of contacting cells of said individual with a multifunctional molecular complex of the present invention, which includes said nucleic acid composition. Here again, the nucleic acid molecule comprises a nucleotide sequence that either encodes a desired peptide or protein, or serves as a  
10 template for functional nucleic acid molecules. The nucleic acid molecule is administered free from retroviral particles. The desired protein may either be a protein which functions within the individual or serves to initiate an immune response.

15        The nucleic acid molecule may be administered to the cells of said individual on either an *in vivo* or *ex vivo* basis, *i.e.*, the contact with the cells of the individual may take place within the body of the individual in accordance with the procedures which are most typically employed, or the contact with the cells of the individual may take place outside the body of the individual by withdrawing cells which it is desired to treat from the body of the individual by various suitable means, followed by contacting of said cells with said nucleic acid molecule, followed in turn by return of said cells to the body of said individual.  
20

25        The method of transferring a nucleic acid composition to the cells of an individual provided by the present invention, includes particularly a method of immunizing an individual against a pathogen. In this method, the nucleic acid composition administered to said cells, comprises a nucleotide sequence that encodes a peptide which comprises at least an epitope identical to, or substantially similar to an epitope displayed on said pathogen as antigen, and said nucleotide sequence is operatively linked to regulatory sequences. The nucleic acid molecule must, of course, be capable of being expressed in the cells of the individual.

30        The method of transferring a nucleic acid composition to the cells of an individual provided by the present invention, further includes methods of immunizing an individual against a hyperproliferative disease or an autoimmune disease. In such methods, the nucleic acid composition administered to the cells of the individual includes a nucleotide sequence that encodes a peptide that comprises at least an epitope identical to or substantially similar to an epitope displayed on a hyperproliferative disease-associated

protein or an autoimmune disease-associated protein, respectively, and is operatively linked to regulatory sequences. Here again, the nucleic acid molecule must be capable of being expressed in the cells of the individual.

The subject may be any mammal, particularly a mammal having  
5 symptoms of a genetically-based disorder. Thus, the subject application finds use in domestic animals, feed stock, such as bovine, ovine, and porcine, as well as primates, particularly humans. The mammalian host may be pregnant, and the intended recipient of the gene-based therapy may be either the gravid female or the fetus or both. In the method of the invention, transfection *in vivo* is obtained by introducing a therapeutic  
10 transcription or expression vector into the mammalian host, either as naked DNA or complexed to lipid carriers, particularly cationic lipid carriers. The constructs may provide for integration into the host cell genome for stable maintenance of the transgene or for episomal expression of the transgene. The introduction into the mammalian host may be by any of several routes, including intravenous or intraperitoneal injection, intratracheally, intrathecally, parenterally, intraarticularly, intramuscularly, etc. Of particular interest is the introduction of a therapeutic expression vector into a circulating bodily fluid. Thus, iv administration and intrathecal administration are of particular interest since the vector may be widely disseminated following such a route of administration.  
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The amount of the compositions of the invention used will be sufficient to provide for adequate dissemination to a variety of tissues after entry of the DNA or complexes into the bloodstream and to provide for a therapeutic level of expression in transfected tissues. A therapeutic level of expression is a sufficient amount of expression to, prevent, treat or palliate a disease of the host mammal. In addition, the dose of the  
25 plasmid DNA expression vector used must be sufficient to produce significant levels of transgene expression in multiple tissues *in vivo* for example, 1 mg of an expression plasmid alone is injected into a mouse to achieve high level expression of the CAT gene in multiple tissues. Other DNA sequences, such as adenovirus VA genes can be included in the administration medium and be co-transfected with the gene of interest. The  
30 presence of genes coding for the adenovirus VA gene product may significantly enhance the translation of mRNA transcribed from the plasmid.

The level and tissues of expression of the recombinant gene may be determined at the mRNA level and/or at the level of polypeptide or protein. Gene product may be quantitated by measuring its biological activity in tissues. For example,

enzymatic activity can be measured by biological assay or by identifying the gene product in transfected cells by immunostaining techniques such as probing with an antibody which specifically recognizes the gene product or a reporter gene product present in the expression cassette. Alternatively, potential therapeutic effects of the gene product can be measured, for example where the DNA sequence of interest encodes GM-CSF, by determining the effects of gene expression on survival of lethally irradiated animals in which the GM-CSF transgene is expressed. Production of significant amounts of a transgene product will substantially prolong the survival of these mice.

Where expression of the polypeptide/protein or even the mRNA itself confers a changed biochemical phenotype upon the host, the presence of a new phenotype or absence of an old phenotype may be evaluated; for example, as a result of transfection of the host cells, there may be enhanced production of pre-existing desirable products formerly produced in insufficient quantities or there may be reduction or even suppression of an undesirable gene product using antisense, ribozyme or co-suppression technologies; in the case of suppression, a reduction of the gene product may be determined. Typically, the therapeutic cassette is not integrated into the host cell genome. If necessary, the treatment can be repeated on an ad hoc basis depending upon the results achieved. If the treatment is repeated, the mammalian host can be monitored to ensure that there is no adverse immune response to the treatment.

The subject compositions can be provided for use in one or more procedures. Kits will usually include the DNA either as naked DNA bearing the reactive linker group or already tethered to carriers. Additionally, lipid carriers may be provided in a separate container for complexing with the provided DNA. The DNA either for direct injection or for complexing with lipid carriers, or the lipid carrier/DNA complexes may be present as concentrates which may be further diluted prior to use or they may be provided at the concentration of use, where the vials may include one or more dosages. Conveniently, single dosages may be provided in syringes, contained in sterilized containers, so that the physicians or veterinarian may employ the syringes directly, where the syringes will have the desired amount and concentration of agents. Thus, the kit may have a plurality of syringes containing the DNA or the DNA/lipid carrier complexes in appropriate proportional amounts. When the syringes contain the formulation for direct use, usually there will be no need for other reagents for use with the method.

The invention finds use in *in vivo* prevention, treatment and/or palliation of a number of diseases. *In vivo* replacement of a gene can be accomplished by techniques

such as homologous recombination or initial knockout of the aberrant gene and subsequent replacement with the desired transgene.

Thus, in accordance with the present invention there is provided a method for the transfer of a nucleic acid composition to target cells on an *in vitro* basis. In this 5 method target cells are contacted with a multifunctional molecular complex which includes said nucleic acid composition. In one embodiment, the target cells have been isolated from an individual, and all of the cells are thus of the same type, and it is not necessary, therefore, for the complex to include a receptor specific binding component. An especially preferred embodiment is one in which a microorganism culture is 10 maintained under fermentation conditions, and a protein product is expressed by the microorganism as a result of the transfer thereto of nucleic acid compositions using the multifunctional molecular complex of the present invention. The protein product is isolated and purified. Here again, a single type of target cell is involved, so that it is not necessary that a receptor specific binding component be present.

15 This method provides for transfer to target cells of a nucleic acid molecule that comprises a nucleotide sequence that either encodes a desired peptide or protein, or serves as a template for functional nucleic acid molecules. The desired protein or functional nucleic acid molecule may be any product of industrial, commercial or scientific interest, *e.g.*, therapeutic agents including vaccines; foodstuffs and nutritional 20 supplements; compounds of agricultural significance such as herbicides and plant growth regulators, insecticides, miticides, rodenticides, and fungicides; compounds useful in animal health such as parasiticides including nematocides; and so forth. The target cells are typically cultures of host cells comprising microorganism cells such as bacteria and yeast, but may also include plant and mammalian cells. The cell cultures are maintained 25 in accordance with fermentation techniques well known in the art, which maximize production of the desired protein or functional nucleic acid molecule, and the fermentation products are harvested and purified by known methods.

It is understood that the examples and embodiments described herein are 30 for illustrative purposes only and that various modifications or changes in light thereof will be suggested to persons skilled in the art and are to be included within the spirit and purview of this application and are considered within the scope of the appended claims. All publications, patents, and patent applications cited herein are hereby incorporated by reference in their entirety for all purposes.

**WHAT IS CLAIMED IS:**

1           1. A substantially water-soluble polymer comprising a first subunit  
2 comprising a first nucleic acid, wherein said first subunit is incorporated into said  
3 polymer using a first subunit precursor comprising said first nucleic acid and an ethylene-  
4 containing moiety.

1           2. The polymer according to claim 1, wherein said first subunit is  
2 covalently attached to a second subunit.

1           3. The polymer according to claim 1, further comprising a cleavable  
2 moiety.

1           4. The polymer according to claim 3, wherein said cleavable moiety  
2 is located between said first subunit and said second subunit.

1           5. The polymer according to claim 3, wherein said cleavable moiety  
2 is a member selected from groups cleaved by change in pH, enzymatic action, reduction,  
3 oxidation, light, heat and combinations thereof.

1           6. The polymer according to claim 5, wherein said cleavable moiety  
2 is cleaved by a process occurring in a biological system.

1           7. The polymer according to claim 6, wherein said cleavable moiety  
2 is a member selected from disulfides, esters, phosphodiesters and combinations thereof.

1           8. The polymer according to claim 1, wherein said first subunit  
2 further comprises a linker group adjoining said first nucleic acid and said ethylene-  
3 containing moiety.

1           9. The polymer according to claim 8, wherein said linker arm  
2 comprises a cleavable moiety.

1           10. The polymer according to claim 8, wherein said cleavable moiety  
2 is a member selected from groups cleaved by change in pH, enzymatic action, reduction,  
3 oxidation, light, heat and combinations thereof.

1           11. The polymer according to claim 9, wherein said cleavable moiety  
2 is cleaved by a process occurring in a biological system.

1           12. The polymer according to claim 10, wherein said cleavable moiety  
2 is a member selected from disulfides, esters, phosphodiesters and combinations thereof.

1           13. The polymer according to claim 1, wherein said ethylene-  
2 containing moiety comprises a member selected from  $-\text{CH}_2=\text{CHX}^1$ ,  $-\text{CH}_2=\text{CX}^2\text{Y}^1$  and  
3 combinations thereof, wherein

4            $\text{X}^1$ ,  $\text{X}^2$  and  $\text{Y}^1$  are members independently selected from H, (=O),  $-\text{NR}^1\text{R}^2$ ,  
5           —OH, and  $-\text{OR}^3$ , wherein

6            $\text{R}^1$ ,  $\text{R}^2$  and  $\text{R}^3$  are members independently selected from H, alkyl,  
7           substituted alkyl, aryl and substituted aryl.

1           14. The polymer according to claim 13, wherein  $\text{R}^1$ ,  $\text{R}^2$  and  $\text{R}^3$  are  
2 independently selected from H, alkyl and substituted alkyl.

1           15. The polymer according to claim 14, wherein  $\text{R}^1$ ,  $\text{R}^2$  and  $\text{R}^3$  are  
2 independently selected from H, alkyl and alkyl substituted with at least one moiety  
3 selected from —OH, —O— and combinations thereof.

1           16. The polymer according to claim 15, wherein at least one of  $\text{R}^1$ ,  $\text{R}^2$   
2 and  $\text{R}^3$  comprises a moiety selected from poly(ethyleneglycol), poly(propyleneglycol) and  
3 combinations thereof.

1           17. The polymer according to claim 1, wherein said polymer comprises  
2 a member selected from acrylate, acrylamide, C<sub>1</sub>-C<sub>6</sub> alkylacrylate, (alkyl)acrylamide,  
3 methylmethacrylate, triethyleneglycolmethacrylate, poly(ethyleneglycol)methacrylate,  
4 hydroxyethylmethacrylate, glycerylmethacrylate, vinyl alcohol, ethylcyanoacrylate and  
5 combinations thereof.

1           18. The polymer according to claim 1, further comprising a tissue-  
2 specific targeting moiety.

1           19. The polymer according to claim 1, further comprising a moiety that  
2 enhances cellular uptake.

1                   **20.**       The polymer according to claim 1, further comprising a nucleic  
2 acid compacting moiety.

1                   **21.**       The polymer according to claim 1, wherein said first nucleic acid is  
2 hybridized to a second nucleic acid.

1                   **22.**       The polymer according to claim 21, wherein said first nucleic acid  
2 is a single-stranded nucleic acid.

1                   **23.**       The polymer according to claim 21, wherein said first nucleic acid  
2 is a double-stranded nucleic acid.

1                   **24.**       The polymer according to claim 22, wherein said second nucleic  
2 acid is a double-stranded nucleic acid.

1                   **25.**       The polymer according to claim 23, wherein said second nucleic  
2 acid is a single-stranded nucleic acid.

1                   **26.**       The polymer according to claim 1, wherein said polymer is a  
2 homopolymer of said first subunit.

1                   **27.**       The polymer according to claim 1, wherein said polymer is a  
2 copolymer of said first subunit and a second subunit.

1                   **28.**       The polymer according to claim 27, wherein said second subunit  
2 comprises a third nucleic acid.

1                   **29.**       The polymer according to claim 28, wherein said third nucleic acid  
2 has a sequence different from that of said first nucleic acid.

1                   **30.**       A polymeric particle comprising a first subunit comprising a first  
2 nucleic acid, wherein said first subunit is incorporated into said polymer using a first  
3 subunit precursor comprising an ethylene-containing moiety.

1                   **31.**       The particle according to claim 30, wherein said first subunit  
2 further comprises a linker group adjoining said first nucleic acid and said ethylene-  
3 containing moiety.

1                   **32.**     The particle according to claim 31, wherein said linker arm  
2 comprises a cleavable moiety.

1                   **33.**     The particle according to claim 31, wherein said cleavable moiety  
2 is a member selected from groups cleaved by change in pH, enzymatic action, reduction,  
3 oxidation, light, heat and combinations thereof.

1                   **34.**     The particle according to claim 32, wherein said cleavable moiety  
2 is cleaved by a process occurring in a biological system.

1                   **35.**     The particle according to claim 33, wherein said cleavable moiety  
2 is a member selected from disulfides, esters and combinations thereof.

1                   **36.**     The particle according to claim 30, wherein said ethylene-  
2 containing moiety comprises a member selected from  $-\text{CH}_2=\text{CHX}^1$ ,  $-\text{CH}_2=\text{CX}^2\text{Y}^1$  and  
3 combinations thereof, wherein

4                   **X**<sup>1</sup>, **X**<sup>2</sup> and **Y**<sup>1</sup> are members independently selected from H, (=O),  $-\text{NR}^1\text{R}^2$ ,  
5                   —OH, and —OR<sup>3</sup>, wherein

6                   **R**<sup>1</sup>, **R**<sup>2</sup> and **R**<sup>3</sup> are members independently selected from H, alkyl,  
7                   substituted alkyl, aryl and substituted aryl.

1                   **37.**     The particle according to claim 36, wherein R<sup>1</sup>, R<sup>2</sup> and R<sup>3</sup> are  
2 independently selected from H, alkyl and substituted alkyl.

1                   **38.**     The particle according to claim 37, wherein R<sup>1</sup>, R<sup>2</sup> and R<sup>3</sup> are  
2 independently selected from H, alkyl and alkyl substituted with at least one moiety  
3 selected from —OH, —O— and combinations thereof.

1                   **39.**     The particle according to claim 38, wherein at least one of R<sup>1</sup>, R<sup>2</sup>  
2 and R<sup>3</sup> comprises a moiety selected from poly(ethyleneglycol), poly(propyleneglycol) and  
3 combinations thereof.

1                   **40.**     The particle according to claim 30, wherein said polymer  
2 comprises a member selected from acrylate, acrylamide, C<sub>1</sub>-C<sub>6</sub> alkylacrylate,  
3 (alkyl)acrylamide, methylmethacrylate, triethyleneglycolmethacrylate,

4 poly(ethyleneglycol)methacrylate, hydroxyethylmethacrylate, glycerylmethacrylate, vinyl  
5 alcohol, ethylcyanoacrylate and combinations thereof.

1           **41.**       The particle according to claim 30, further comprising a tissue-  
2 specific targeting moiety.

1           **42.**       The particle according to claim 30, further comprising a moiety  
2 that enhances cellular uptake.

1           **43.**       The particle according to claim 30, further comprising a nucleic  
2 acid compacting moiety.

1           **44.**       The particle according to claim 30, wherein said first nucleic acid  
2 is hybridized to a second nucleic acid.

1           **45.**       The particle according to claim 44, wherein said first nucleic acid  
2 is a single-stranded nucleic acid.

1           **46.**       The particle according to claim 44, wherein said first nucleic acid  
2 is a double-stranded nucleic acid.

1           **47.**       The particle according to claim 45, wherein said second nucleic  
2 acid is a double-stranded nucleic acid.

1           **48.**       The particle according to claim 46, wherein said second nucleic  
2 acid is a single-stranded nucleic acid.

1           **49.**       The particle according to claim 30, wherein said polymer is a  
2 homopolymer of said first subunit.

1           **50.**       The particle according to claim 30, wherein said polymer is a  
2 copolymer of said first subunit and a second subunit.

1           **51.**       The particle according to claim 50, wherein said second subunit  
2 comprises a third nucleic acid.

1           **52.**       The particle according to claim 51, wherein said third nucleic acid  
2 has a sequence different from that of said first nucleic acid.

1                   **53.**       The particle according to claim 30, wherein said particle is  
2 substantially water-soluble.

1                   **54.**       The particle according to claim 30, wherein said particle is  
2 substantially water-insoluble.

1                   **55.**       The particle according to claim 30, further comprising a bioactive  
2 compound encapsulated by said polymer.

1                   **56.**       A pharmaceutical formulation comprising a pharmaceutically  
2 acceptable carrier and a substantially water-soluble polymer comprising a first subunit  
3 comprising a first nucleic acid, wherein said first subunit is incorporated into said  
4 polymer using a first subunit precursor comprising said first nucleic acids and an  
5 ethylene-containing moiety.

1                   **57.**       A pharmaceutical formulation comprising a pharmaceutically  
2 acceptable carrier and a polymeric particle comprising a first subunit comprising a first  
3 nucleic acid, wherein said first subunit is incorporated into said polymer using a first  
4 subunit precursor comprising said first nucleic acid and an ethylene-containing moiety.

1                   **58.**       A method for treating or preventing a condition, the method  
2 comprising administering to a subject a substantially water-soluble polymer in an amount  
3 effective to treat or prevent said condition, said polymer comprising a first subunit  
4 comprising a first nucleic acid, wherein said first subunit is incorporated into said  
5 polymer using a first subunit precursor comprising said first nucleic acid and an ethylene-  
6 containing moiety.

1                   **59.**       A method for treating or preventing a condition, the method  
2 comprising administering to a subject a polymeric particle in an amount effective to treat  
3 or prevent said condition, said particle comprising a first subunit comprising a first  
4 nucleic acid, wherein said first subunit is incorporated into said polymer using a first  
5 subunit precursor comprising said first nucleic acid and an ethylene-containing moiety.

1                   **60.**       A method for introducing a polynucleotide into a eukaryotic cell in  
2 a living animal comprising contacting the cell with a composition comprising:

3           a substantially water-soluble polymer comprising a first subunit  
4   comprising a first nucleic acid, wherein said first subunit is incorporated into said  
5   polymer using a first subunit precursor comprising said first nucleic acid and an ethylene-  
6   containing moiety.

1           **61.**   The method of claim **60**, wherein said composition is administered  
2   in an amount comprising about 0.5 µg to 20 mg of nucleic acid.

1           **62.**   The method of claim **60**, wherein the eukaryotic cell is a  
2   mammalian cell.

1           **63.**   The method of claim **60**, wherein the composition is administered  
2   by a route selected from the group consisting of oral, transdermal, systemic and inhalation  
3   routes.

1           **64.**   The method of claim **63** wherein the composition is administered  
2   transdermally by high velocity impaction administration to the skin surface.

1           **65.**   A method for introducing a polynucleotide into a eukaryotic cell in  
2   a living animal comprising contacting the cell with a composition comprising:  
3                a polymeric particle comprising a first subunit comprising a first nucleic  
4   acid, wherein said first subunit is incorporated into said polymer using a first subunit  
5   precursor comprising said nucleic acid and an ethylene-containing moiety.

## DECLARATION (37 CFR 1.63) FOR UTILITY OR DESIGN APPLICATION USING AN APPLICATION DATA SHEET (37 CFR 1.76)

As the below named inventor(s), I/we declare that:

This declaration is directed to:

The attached application, or  
 Application No. 09/857,378, filed on June 4, 2001,  
 as amended on \_\_\_\_\_ (if applicable);

I/we believe that I/we am/are the original and first inventor(s) of the subject matter which is claimed and for which a patent is sought;

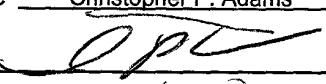
I/we have reviewed and understand the contents of the above-identified application, including the claims, as amended by any amendment specifically referred to above;

I/we acknowledge the duty to disclose to the United States Patent and Trademark Office all information known to me/us to be material to patentability as defined in 37 CFR 1.56, including material information which became available between the filing date of the prior application and the National or PCT International filing date of the continuation-in-part application, if applicable; and

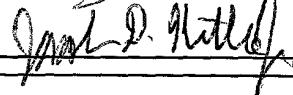
All statements made herein of my/our own knowledge are true, all statements made herein on information and belief are believed to be true, and further that these statements were made with the knowledge that willful false statements and the like are punishable by fine or imprisonment, or both, under 18 U.S.C. 1001, and may jeopardize the validity of the application or any patent issuing thereon.

### FULL NAME OF INVENTOR(S)

Inventor one Christopher P. Adams Date: 11/6/01

Signature:  Citizen of: United States

Inventor two Joseph D. Kittle, Jr. Date: Sept 10, 2001

Signature:  Citizen of: United States

Inventor three \_\_\_\_\_ Date: \_\_\_\_\_

Signature: \_\_\_\_\_ Citizen of: \_\_\_\_\_

Inventor four \_\_\_\_\_ Date: \_\_\_\_\_

Signature: \_\_\_\_\_ Citizen of: \_\_\_\_\_

Additional inventors are being named on \_\_\_\_\_ additional form(s) attached hereto.

Burden Hour Statement: This collection of information is required by 35 U.S.C. 115 and 37 CFR 1.63. The information is used by the public to file (and the PTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This form is estimated to take 1 minute to complete. This time will vary depending upon the needs of the individual case. Any comments on the amount of time you are required to complete this form should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, Washington, DC 20231. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Assistant Commissioner for Patents, Washington, DC 20231.

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**POWER OF ATTORNEY OR  
AUTHORIZATION OF AGENT**

<b>Application Number</b>	09/857,378
<b>Filing Date</b>	June 4, 2001
<b>First Named Inventor</b>	Adams, Christopher P.
<b>Title</b>	Method For The Immobilization Of Oligonucleotides
<b>Group Art Unit</b>	Not Yet Assigned
<b>Examiner Name</b>	Not Yet Assigned
<b>Attorney Docket Number</b>	18422-000310US

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I am the:

Applicant/Inventor.

Assignee of record of the entire interest. See 37 CFR 3.71.

Statement under 37 CFR 3.73(b) is enclosed. (Form PTO/SB/96).

**SIGNATURE of Applicant or Assignee of Record**

Name      Joseph D. Kittle, Jr.

Signature

Date

9/10/2001

NOTE: Signatures of all the inventors or assignees of record of the entire interest or their representative(s) are required.  
Submit multiple forms if more than one signature is required, see below\*.

\*Total of \_\_\_\_\_ forms are submitted.

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<b>POWER OF ATTORNEY OR AUTHORIZATION OF AGENT</b>	Application Number	08/857,378
	Filing Date	June 4, 2001
	First Named Inventor	Adams, Christopher P.
	Title	Method For The Immobilization Of Oligonucleotides
	Group Art Unit	Not Yet Assigned
	Examiner Name	Not Yet Assigned
	Attorney Docket Number	18422-000310US

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Telephone	Fax		

I am the:

Applicant/inventor,

Assignee of record of the entire interest. See 37 CFR 3.71.

*Statement under 37 CFR 3.73(b) is enclosed. (Form PTO/SB/96).*

**SIGNATURE of Applicant or Assignee of Record**

Name	Christopher P. Adams
Signature	
Date	11/15/01

NOTE: Signatures of all the inventors or assignees of record of the entire interest or their representative(s) are required.  
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SF 1201454 v1

**VERIFIED STATEMENT (DECLARATION) CLAIMING SMALL ENTITY STATUS  
(37 CFR 1.9(f) & 1.27(b)) - INDEPENDENT INVENTOR**

Applicant or Patentee:  
Application or Patent No.:  
Filed or Issued:  
Title:

Christopher P. Adams, et al.

09/857,378

June 4, 2001

METHOD FOR THE IMMOBILIZATION OF OLIGONUCLEOTIDES

As a below named inventor, I hereby declare that I qualify as an independent inventor as defined in 37 CFR 1.9(c) for purposes of paying reduced fees to the Patent and Trademark Office regarding the invention entitled METHOD FOR THE IMMOBILIZATION OF OLIGONUCLEOTIDES described in:

the specification filed herewith;  
 Application No. 09/857,378, filed June 4, 2001;  
 Patent No. \_\_\_\_\_, issued \_\_\_\_\_.

I have not assigned, granted, conveyed or licensed and am under no obligation under contract or law to assign, grant, convey or license, any rights in the invention to any person who would not qualify as an independent inventor under 37 CFR 1.9(c) if that person made the invention, or by any concern that would not qualify as a small business concern under 37 CFR 1.9(d), or a nonprofit organization under 37 CFR 1.9(e).

Each person, concern or organization to which I have assigned, granted, conveyed or licensed or am under an obligation under contract or law to assign, grant, convey or license any rights in the invention is listed below.\*

No such person, concern or organization.  
 Persons, concerns or organizations listed below.\*

\*NOTE: Separate verified statements are required from each named person, concern or organization having rights to the invention averring to their status as small entities. (37 CFR 1.27)

Name:  
Address:

\_\_\_\_\_  
\_\_\_\_\_

Individual                     Small Business Concern                     Nonprofit Organization

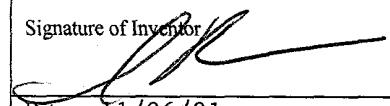
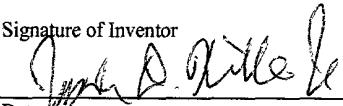
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\_\_\_\_\_

Individual                     Small Business Concern                     Nonprofit Organization

I acknowledge the duty to file, in this application or patent, notification of any change in status resulting in loss of entitlement to small entity status prior to paying, or at the time of paying, the earliest of the issue fee or any maintenance fee due after the date on which status as a small entity is no longer appropriate. (37 CFR 1.28(b))

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application, any patent issuing thereon, or any patent to which this verified statement is directed.

Name of Inventor	Name of Inventor	Name of Inventor
Christopher P. Adams	Joseph D. Kittle, Jr.	
Signature of Inventor 	Signature of Inventor 	Signature of Inventor
Date 11/06/01	Date	Date Sept 10/2001

SF 1251473 v1